

Peridynamic States

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Objective of peridynamics

Some limitations of the standard theory of solid mechanics

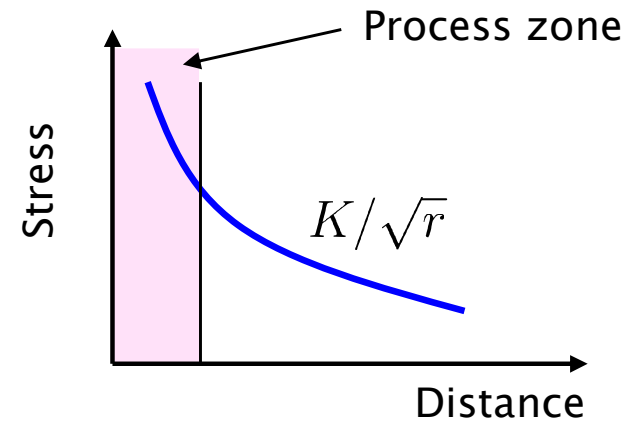
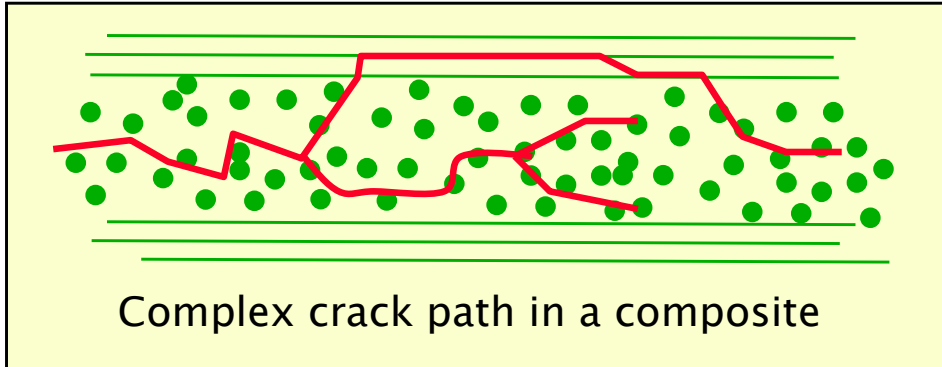
- It is incompatible with the essential physical nature of particles and cracks.
 - Can't apply the PDEs directly.
- Can't easily include long-range interactions.

What the peridynamic theory seeks to do

- To predict the mechanics of continuous and discontinuous media with **mathematical consistency**.
 - Everything should emerge from the same continuum model.

Cracks vs. continua: Why this issue is important

- Typical approaches require some fix at the discretized level.
- LEFM adds extra laws that tell a crack what to do.
 - These laws are known only in idealized cases.



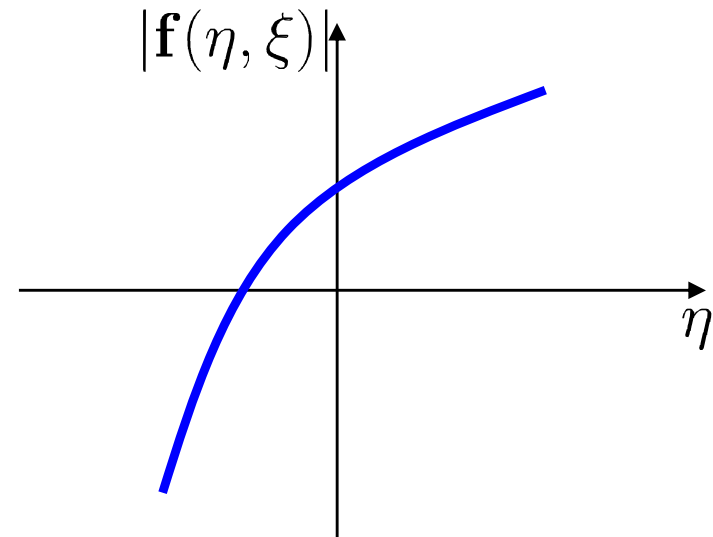
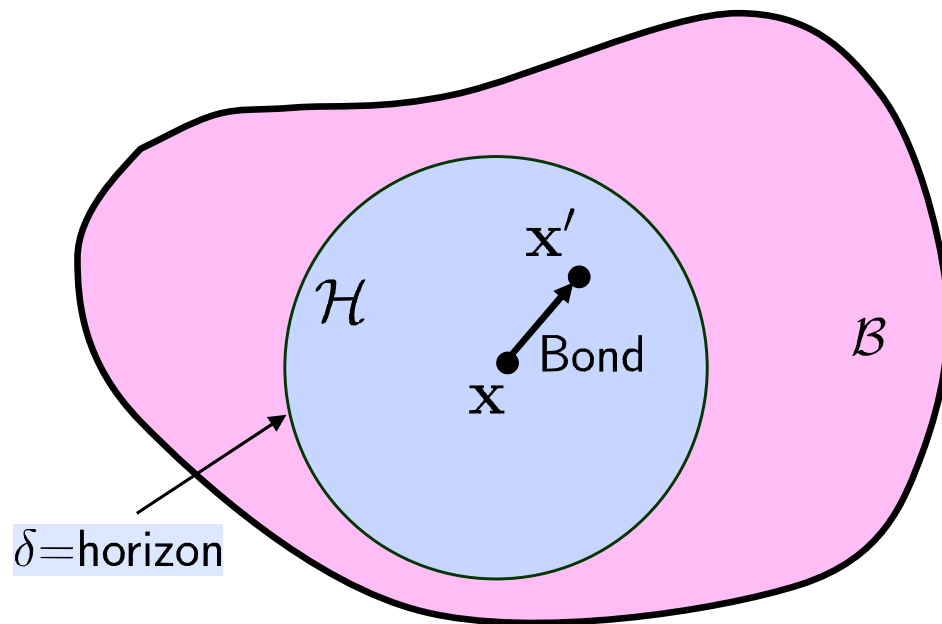
The reality of fracture may be too complex to represent in the form

$$\dot{a} = f(K)$$

Original concept (2000): Continuum as a network of bonds

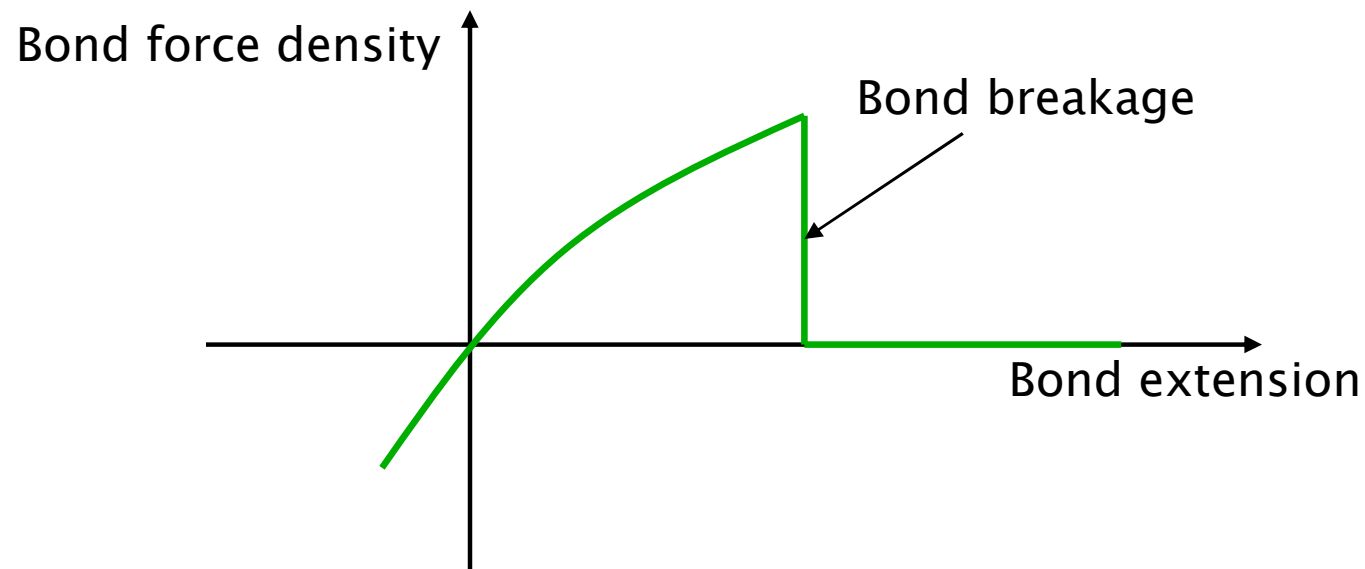
- Any point \mathbf{x} interacts directly with other points within a finite distance δ called the “horizon.” Equation of motion:

$$\rho(\mathbf{x})\ddot{\mathbf{u}}(\mathbf{x}, t) = \int_{\mathcal{H}} \mathbf{f}(\mathbf{u}(\mathbf{x}', t) - \mathbf{u}(\mathbf{x}, t), \mathbf{x}' - \mathbf{x}) dV_{\mathbf{x}'} + \mathbf{b}(\mathbf{x}, t)$$

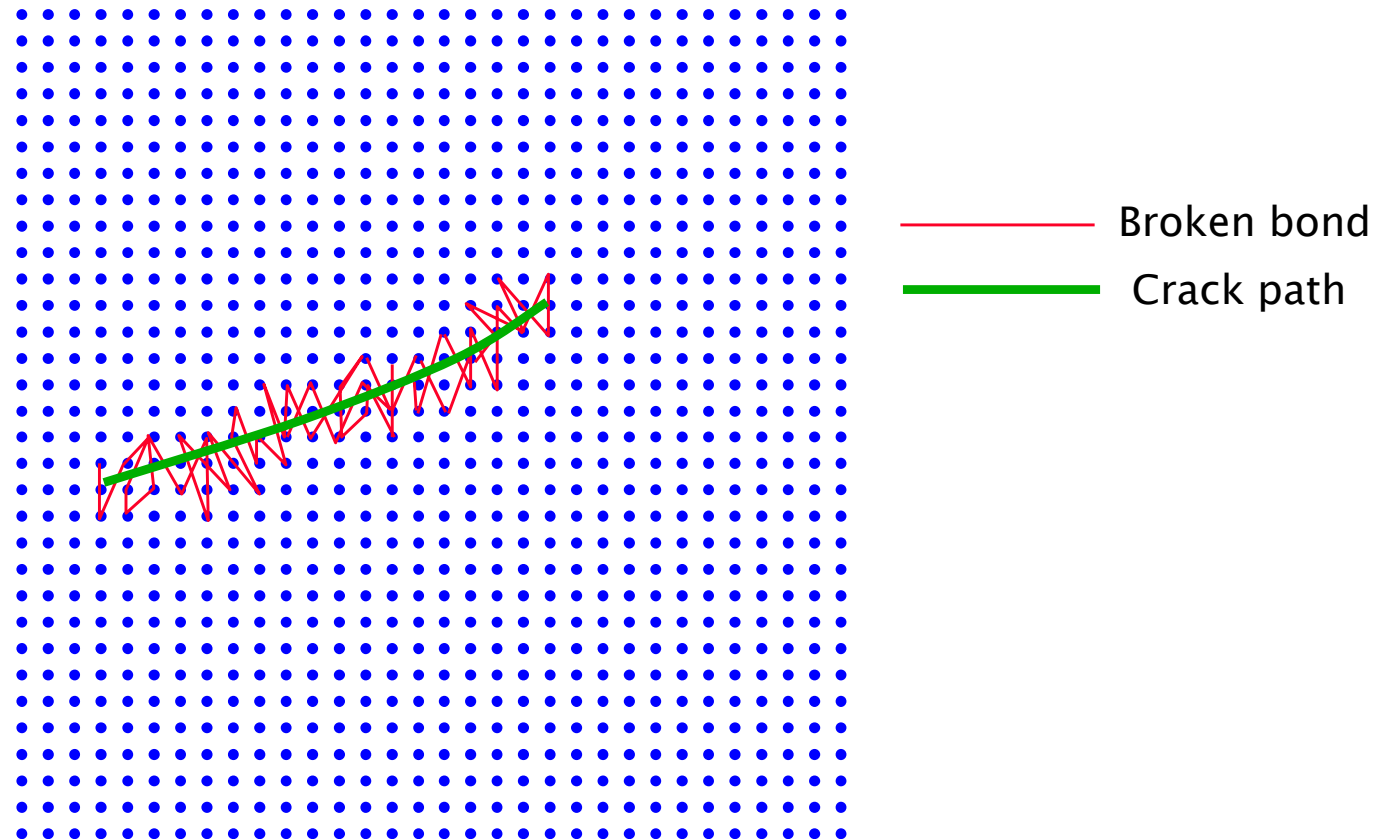


How damage and fracture are modeled

- Bonds can break irreversibly according to some criterion.
- Broken bonds carry no force.



Bond breakage forms cracks “autonomously”

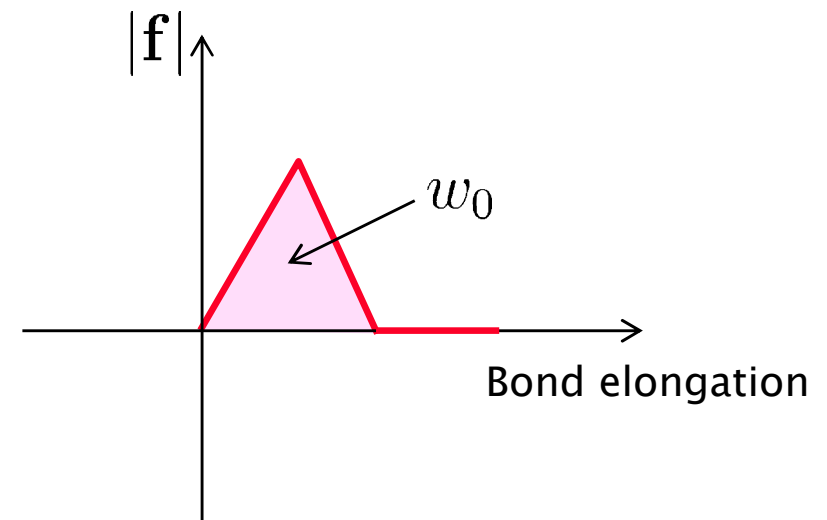
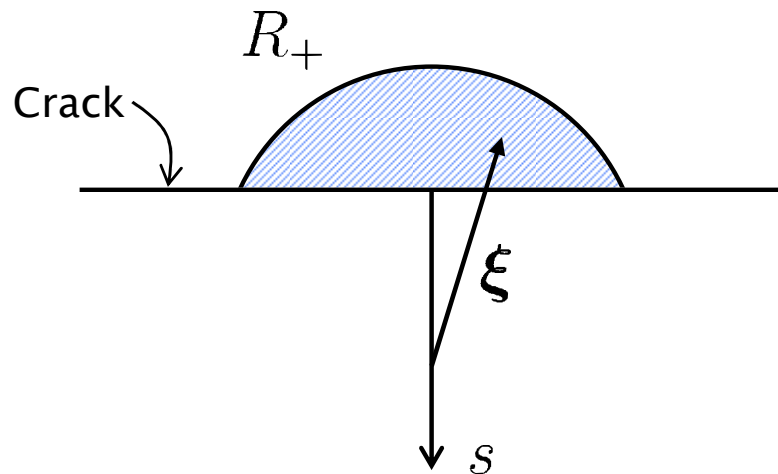


When a bond breaks, its load is shifted to its neighbors, leading to progressive failure.

Energy balance for an advancing crack

If the work required to break the bond ξ is $w_0(\xi)$, then the energy release rate is found by summing this work per unit crack area (J. Foster):

$$G = \int_0^\delta \int_{R_+} w_0(\xi) dV_\xi ds$$



There is also a version of the J-integral that applies in this theory.

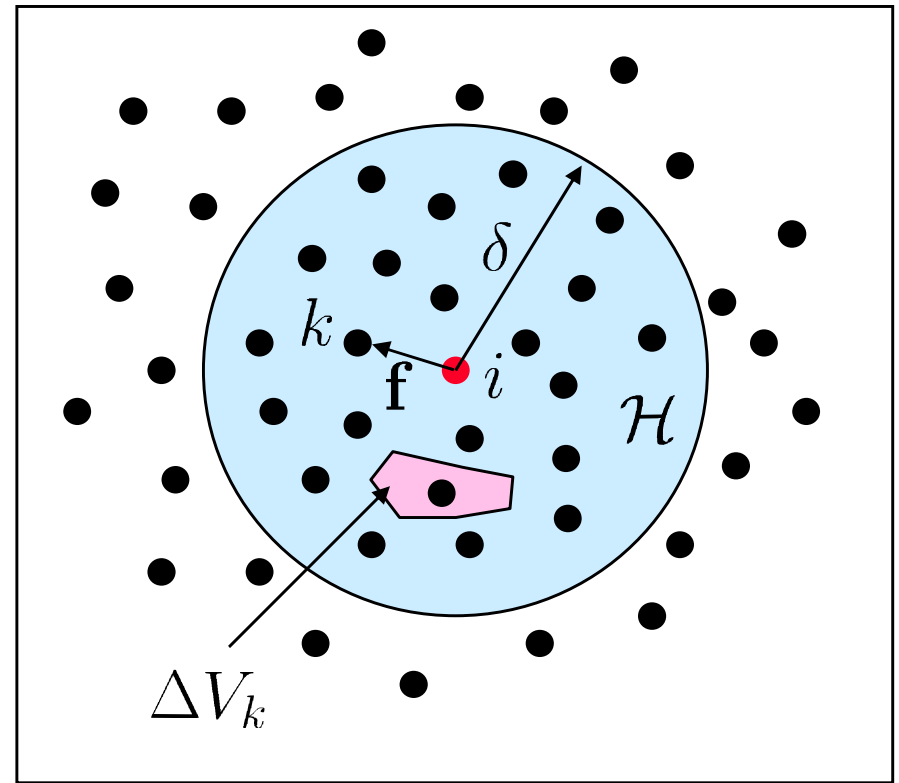
EMU numerical method

- Integral is replaced by a finite sum: resulting method is meshless and Lagrangian.

$$\rho \ddot{\mathbf{y}}(\mathbf{x}, t) = \int_{\mathcal{H}} \mathbf{f}(\mathbf{x}', \mathbf{x}, t) dV_{\mathbf{x}'} + \mathbf{b}(\mathbf{x}, t)$$

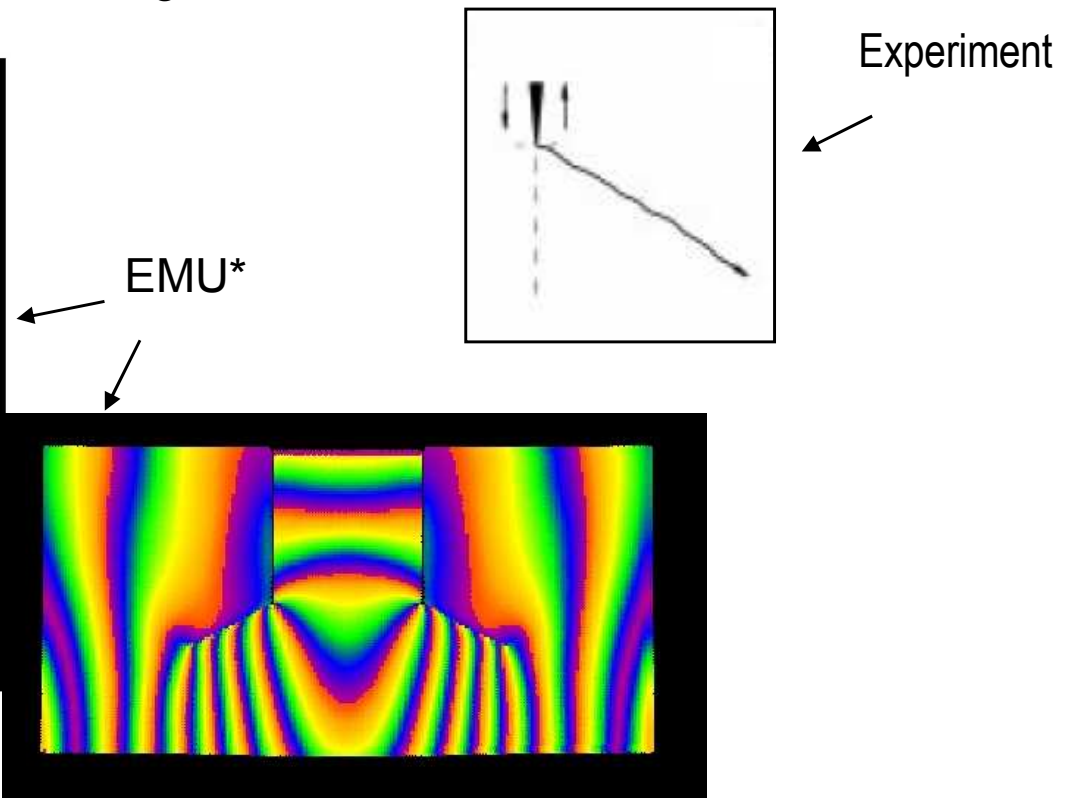
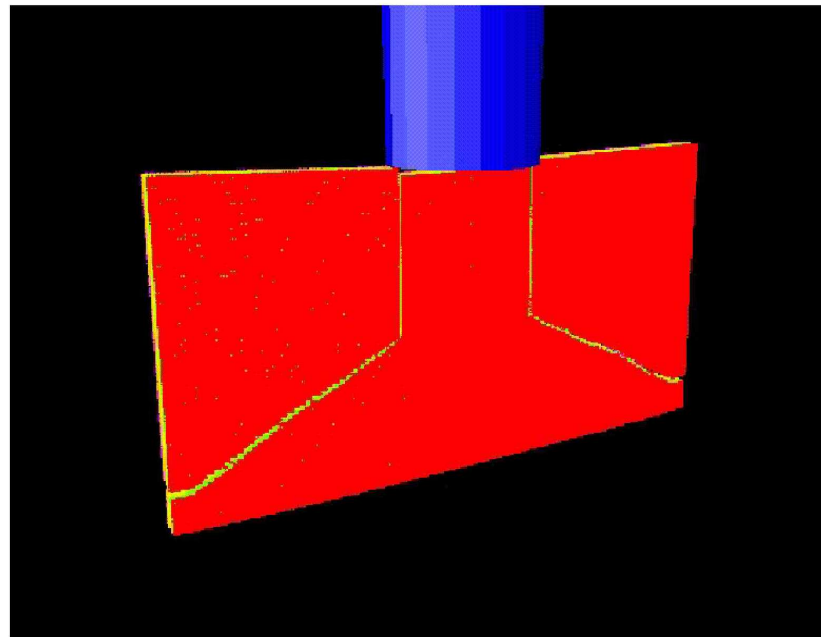


$$\rho \ddot{\mathbf{y}}_i^n = \sum_{k \in \mathcal{H}} \mathbf{f}(\mathbf{x}_k, \mathbf{x}_i, t) \Delta V_k + \mathbf{b}_i^n$$



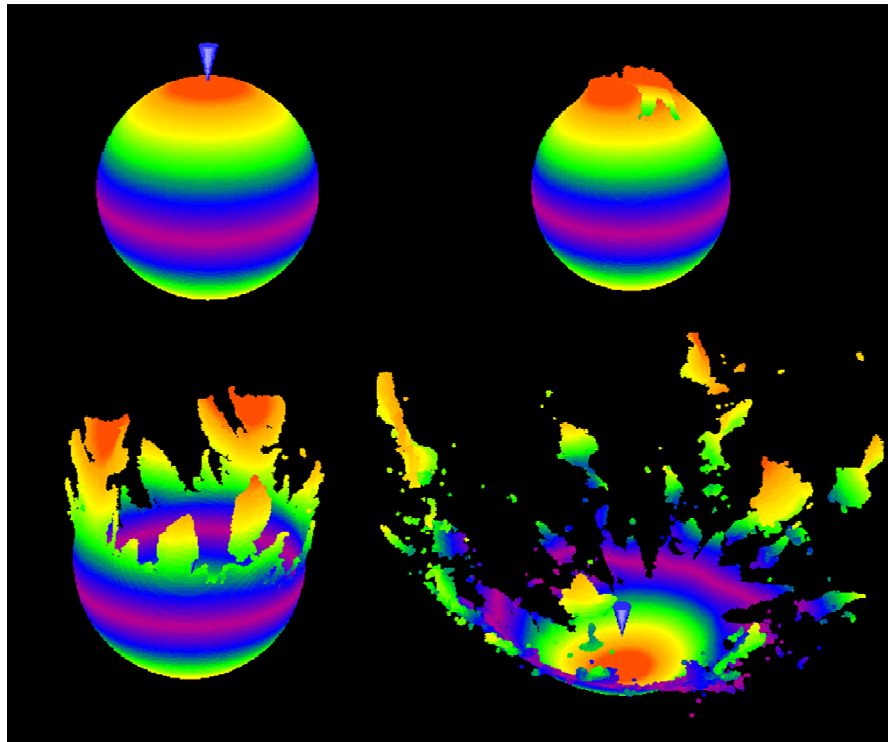
Dynamic fracture in a hard steel plate

- Dynamic fracture in maraging steel (Kalthoff & Winkler, 1988)
 - Mode-II loading at notch tips results in mode-I cracks at 70deg angle.
 - 3D EMU model reproduces the crack angle.



S. A. Silling, Dynamic fracture modeling with a meshfree peridynamic code, in *Computational Fluid and Solid Mechanics 2003*, K.J. Bathe, ed., Elsevier, pp. 641–644.

Dynamic fracture in membranes



EMU model of a balloon penetrated
by a fragment

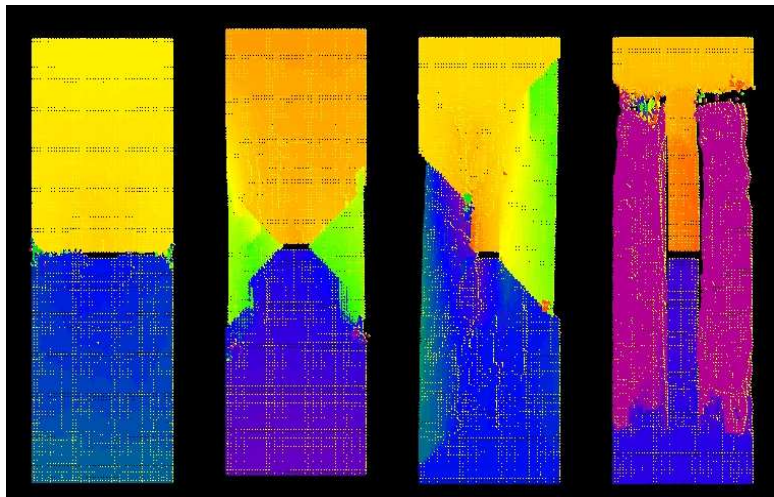


Early high speed photograph by Harold Edgerton
(MIT collection)

<http://mit.edu/6.933/www/Fall2000/edgerton/edgerton.ppt>

Splitting and fracture mode change in composites

- Distribution of fiber directions between plies strongly influences the way cracks grow.



EMU simulations for different layups



Typical crack growth in a notched laminate
(photo courtesy Boeing)



Limitations of the original theory

- Pair interactions imply Poisson ratio = $1/4$.
- Can't use traditional stress-strain models.
- Can't enforce plastic incompressibility.

New approach

- Retain idea of bond forces.
- But bond forces depend on the collective deformation of the family.

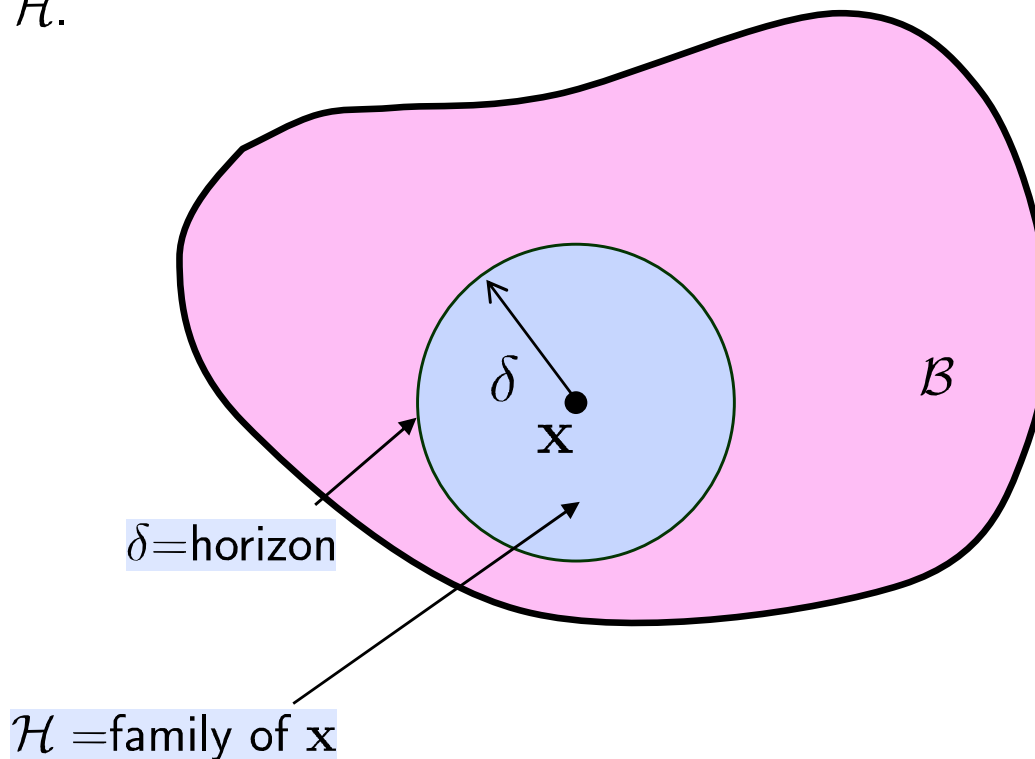


Some references

- S.A. Silling, M. Epton, O. Weckner, J. Xu, and E. Askari, Peridynamic states and constitutive modeling, J. Elast. 88 (2007) 151–184
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- R.B. Lehoucq, S.A. Silling, Force flux and the peridynamic stress tensor, JMPS 56 (2008) 1566–1577
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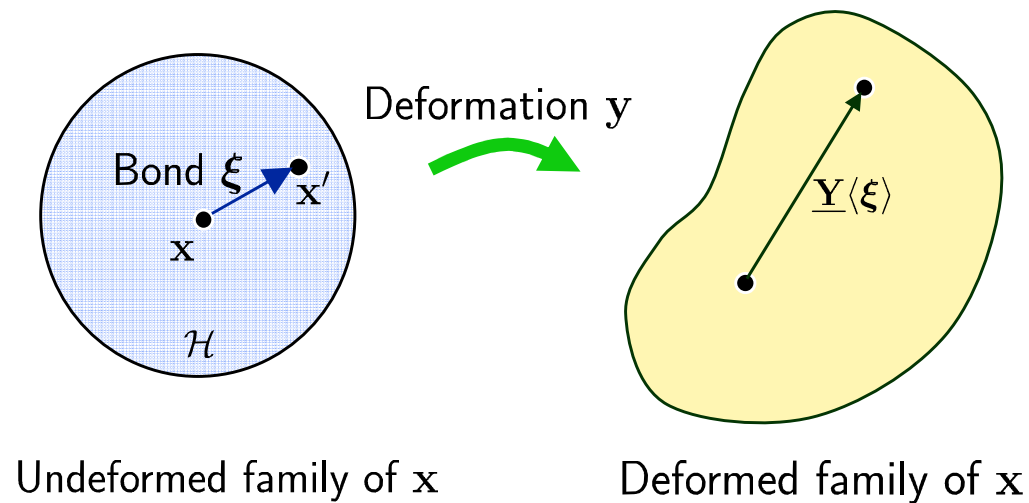
Peridynamics basics: Horizon and family

- Any point \mathbf{x} interacts directly with other points within a finite distance δ called the “horizon.”
- The material within a distance δ of \mathbf{x} is called the “family” of \mathbf{x} , \mathcal{H} .



Why we need states

We want to express the idea that the strain energy density at \mathbf{x} depends **collectively** on the deformation of the family of \mathbf{x} .



Standard:

$$W\left(\frac{\partial \mathbf{y}}{\partial \mathbf{x}}\right)$$

Peridynamic:

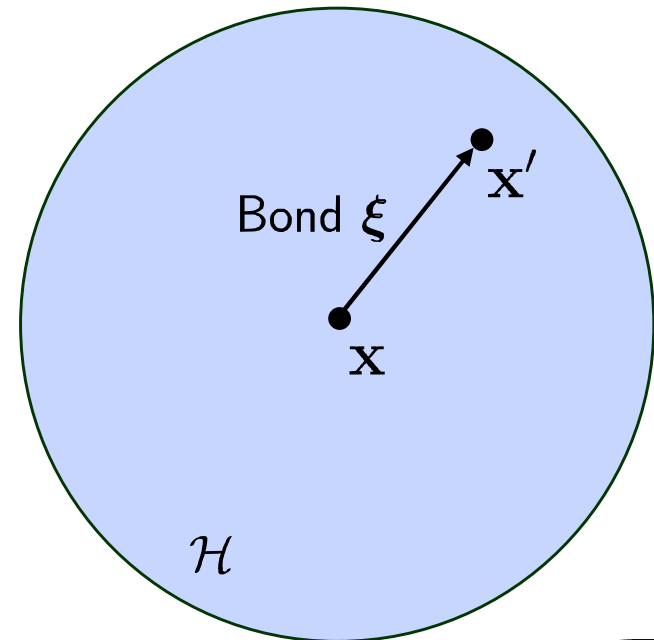
$$W(\underline{\mathbf{Y}})$$

Definition of a state

- A state is a function on \mathcal{H} .
- A vector state \underline{A} maps each bond $\xi \in \mathcal{H}$ to a vector written $\underline{A}\langle\xi\rangle$.
- Scalar states: $\underline{A}\langle\xi\rangle$ is scalar valued.
- Double states map pairs of bonds to second order tensors: $\underline{A}\langle\xi, \zeta\rangle$.

$$\xi = \mathbf{x}' - \mathbf{x}$$

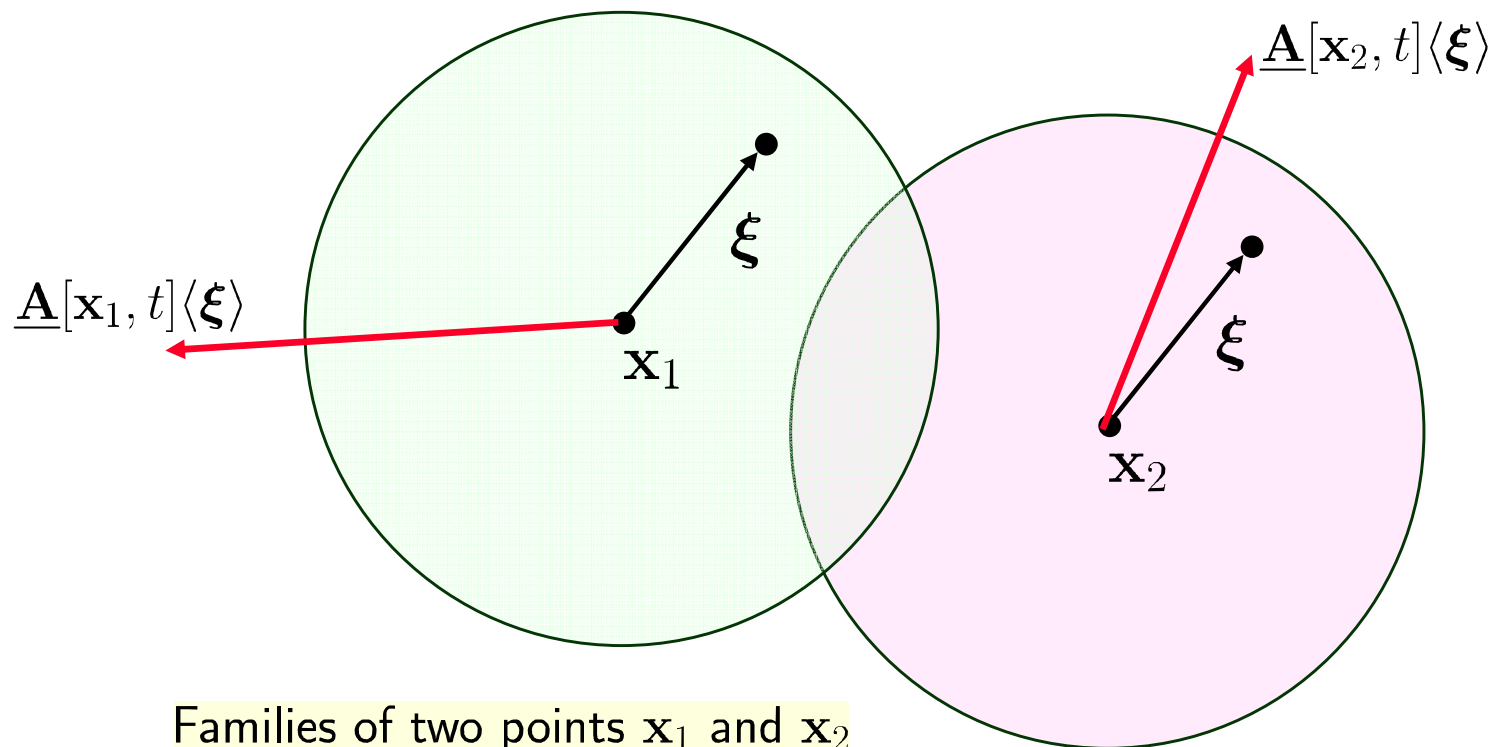
Bonds are defined in the reference configuration.



State fields

- States can depend on position (in the reference configuration) and time.

$$\underline{\mathbf{A}}[\mathbf{x}, t] \langle \xi \rangle$$



Families of two points \mathbf{x}_1 and \mathbf{x}_2



Dot product of two states

- Suppose $\underline{\mathbf{A}}$ and $\underline{\mathbf{B}}$ are vector states.
- Define a scalar called the dot product of $\underline{\mathbf{A}}$ and $\underline{\mathbf{B}}$ by

$$\underline{\mathbf{A}} \bullet \underline{\mathbf{B}} = \int_{\mathcal{H}} \underline{\mathbf{A}}\langle \xi \rangle \cdot \underline{\mathbf{B}}\langle \xi \rangle dV_{\xi}.$$

- In components,

$$\underline{\mathbf{A}} \bullet \underline{\mathbf{B}} = \int_{\mathcal{H}} \underline{A}_i\langle \xi \rangle \underline{B}_i\langle \xi \rangle dV_{\xi}.$$

- Norm of a vector state:

$$||\underline{\mathbf{A}}|| = \sqrt{\underline{\mathbf{A}} \bullet \underline{\mathbf{A}}}$$



Dot product of two states, ctd.

- Suppose \underline{a} and \underline{b} are scalar states.

$$\underline{a} \bullet \underline{b} = \int_{\mathcal{H}} \underline{a} \langle \xi \rangle \underline{b} \langle \xi \rangle dV_{\xi}.$$

- *Point product* is a scalar state:

$$(\underline{ab}) \langle \xi \rangle = \underline{a} \langle \xi \rangle \underline{b} \langle \xi \rangle$$

Functions of states, Frechet derivatives

- Let $\Psi(\underline{\mathbf{A}})$ be a scalar valued function of a vector state.
- How much does Ψ change if we change $\underline{\mathbf{A}}$? Suppose there is a vector state $\Psi_{\underline{\mathbf{A}}}$ such that

$$\Psi(\underline{\mathbf{A}} + \underline{\mathbf{a}}) = \Psi(\underline{\mathbf{A}}) + \Psi_{\underline{\mathbf{A}}} \bullet \underline{\mathbf{a}} + o(||\underline{\mathbf{a}}||)$$

for any vector state $\underline{\mathbf{a}}$.

- $\Psi_{\underline{\mathbf{A}}}(\underline{\mathbf{A}})$ is the Fréchet derivative of Ψ at $\underline{\mathbf{A}}$.

Less than first order.



Maurice Rene Frechet

Concept is similar to the gradient in \mathbb{R}^3 , e.g.,

$$f(\mathbf{x} + \delta\mathbf{x}) = f(\mathbf{x}) + f_{\mathbf{x}}(\mathbf{x}) \cdot \delta\mathbf{x} + o(|\delta\mathbf{x}|)$$

except that $\underline{\mathbf{A}}$ lives in an infinite dimensional space.



Frechet derivative examples: 1

- Find the Frechet derivative of $\Psi(\underline{\mathbf{A}}) = \underline{\mathbf{A}} \bullet \underline{\mathbf{A}}$:

$$\begin{aligned}\Psi(\underline{\mathbf{A}} + \underline{\mathbf{a}}) &= (\underline{\mathbf{A}} + \underline{\mathbf{a}}) \bullet (\underline{\mathbf{A}} + \underline{\mathbf{a}}) \\ &= \int (\underline{\mathbf{A}}\langle\xi\rangle + \underline{\mathbf{a}}\langle\xi\rangle) \cdot (\underline{\mathbf{A}}\langle\xi\rangle + \underline{\mathbf{a}}\langle\xi\rangle) dV_\xi \\ &= \underline{\mathbf{A}} \bullet \underline{\mathbf{A}} + 2 \int \underline{\mathbf{A}}\langle\xi\rangle \cdot \underline{\mathbf{a}}\langle\xi\rangle dV_\xi + O(\|\underline{\mathbf{a}}\|) \\ &= \Psi(\underline{\mathbf{A}}) + 2\underline{\mathbf{A}} \bullet \underline{\mathbf{a}} + O(\|\underline{\mathbf{a}}\|)\end{aligned}$$

Therefore

$$\Psi_{\underline{\mathbf{A}}} = 2\underline{\mathbf{A}}.$$



Frechet derivative examples: 2

- Find the Frechet derivative of $\Psi(\underline{\mathbf{A}}) = \int |\underline{\mathbf{A}}\langle\xi\rangle| dV_\xi$:

$$\begin{aligned}\Psi(\underline{\mathbf{A}} + \underline{\mathbf{a}}) &= \int |\underline{\mathbf{A}}\langle\xi\rangle + \underline{\mathbf{a}}\langle\xi\rangle| dV_\xi \\&= \int \sqrt{(\underline{\mathbf{A}}\langle\xi\rangle + \underline{\mathbf{a}}\langle\xi\rangle) \cdot (\underline{\mathbf{A}}\langle\xi\rangle + \underline{\mathbf{a}}\langle\xi\rangle)} dV_\xi \\&= \int |\underline{\mathbf{A}}\langle\xi\rangle| \sqrt{1 + \frac{2\underline{\mathbf{A}}\langle\xi\rangle \cdot \underline{\mathbf{a}}\langle\xi\rangle}{|\underline{\mathbf{A}}\langle\xi\rangle|^2} + \dots} dV_\xi \\&= \int |\underline{\mathbf{A}}\langle\xi\rangle| \left(1 + \frac{\underline{\mathbf{A}}\langle\xi\rangle \cdot \underline{\mathbf{a}}\langle\xi\rangle}{|\underline{\mathbf{A}}\langle\xi\rangle|^2} + \dots \right) dV_\xi \\&= \Psi(\underline{\mathbf{A}}) + \int \frac{\underline{\mathbf{A}}\langle\xi\rangle \cdot \underline{\mathbf{a}}\langle\xi\rangle}{|\underline{\mathbf{A}}\langle\xi\rangle|} dV_\xi + \dots\end{aligned}$$

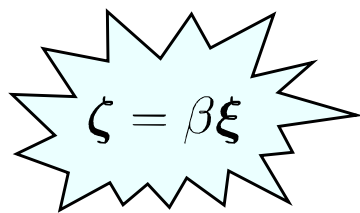
Therefore

$$\Psi_{\underline{\mathbf{A}}} = \frac{\underline{\mathbf{A}}}{|\underline{\mathbf{A}}|}.$$

Frechet derivative examples: 3

- Find the Frechet derivative of $\Psi(\underline{\mathbf{A}}) = \int \underline{\mathbf{A}}\langle \boldsymbol{\xi} \rangle \cdot \underline{\mathbf{A}}\langle \beta \boldsymbol{\xi} \rangle dV_{\boldsymbol{\xi}}$ where β is a constant:

$$\Psi(\underline{\mathbf{A}} + \underline{\mathbf{a}}) = \int (\underline{\mathbf{A}}\langle \boldsymbol{\xi} \rangle + \underline{\mathbf{a}}\langle \boldsymbol{\xi} \rangle) \cdot (\underline{\mathbf{A}}\langle \beta \boldsymbol{\xi} \rangle + \underline{\mathbf{a}}\langle \beta \boldsymbol{\xi} \rangle) dV_{\boldsymbol{\xi}}$$



$$= \Psi(\underline{\mathbf{A}}) + \int (\underline{\mathbf{A}}\langle \boldsymbol{\xi} \rangle \cdot \underline{\mathbf{a}}\langle \beta \boldsymbol{\xi} \rangle + \underline{\mathbf{A}}\langle \beta \boldsymbol{\xi} \rangle \cdot \underline{\mathbf{a}}\langle \boldsymbol{\xi} \rangle) dV_{\boldsymbol{\xi}} + \dots$$

$$= \Psi(\underline{\mathbf{A}}) + \int \underline{\mathbf{A}}\langle \beta^{-1} \boldsymbol{\zeta} \rangle \cdot \underline{\mathbf{a}}\langle \boldsymbol{\zeta} \rangle (\beta^{-3} dV_{\boldsymbol{\zeta}}) + \int \underline{\mathbf{A}}\langle \beta \boldsymbol{\xi} \rangle \cdot \underline{\mathbf{a}}\langle \boldsymbol{\xi} \rangle dV_{\boldsymbol{\xi}} + \dots$$

$$= \Psi(\underline{\mathbf{A}}) + \int (\beta^{-3} \underline{\mathbf{A}}\langle \beta^{-1} \boldsymbol{\xi} \rangle + \underline{\mathbf{A}}\langle \beta \boldsymbol{\xi} \rangle) \cdot \underline{\mathbf{a}}\langle \boldsymbol{\xi} \rangle dV_{\boldsymbol{\xi}} + \dots$$

Therefore

$$\Psi_{\underline{\mathbf{A}}}\langle \boldsymbol{\xi} \rangle = \beta^{-3} \underline{\mathbf{A}}\langle \beta^{-1} \boldsymbol{\xi} \rangle + \underline{\mathbf{A}}\langle \beta \boldsymbol{\xi} \rangle.$$



Frechet derivative examples: 4

- Find the Frechet derivative of $\Psi(\underline{\mathbf{A}}) = \mathbf{c} \cdot \underline{\mathbf{A}}\langle \xi_0 \rangle$ where \mathbf{c} is a constant vector and $\xi_0 \in \mathcal{H}$ is a given bond:

$$\begin{aligned}\Psi(\underline{\mathbf{A}} + \underline{\mathbf{a}}) &= \int \mathbf{c} \cdot (\underline{\mathbf{A}}\langle \xi \rangle + \underline{\mathbf{a}}\langle \xi \rangle) \Delta(\xi - \xi_0) dV_\xi \\ &= \Psi(\underline{\mathbf{A}}) + \int \Delta(\xi - \xi_0) \mathbf{c} \cdot \underline{\mathbf{a}}\langle \xi \rangle dV_\xi\end{aligned}$$

where Δ is the Dirac delta function. Therefore

$$\Psi_{\underline{\mathbf{A}}}\langle \xi \rangle = \Delta(\xi - \xi_0) \mathbf{c}.$$



Frechet derivative examples: 5

- Find the Frechet derivative of $\Psi(\underline{\mathbf{A}}) = \int f(\underline{\mathbf{A}}\langle\xi\rangle) dV_\xi$ where $f(\mathbf{v})$ is a scalar-valued function of a vector:

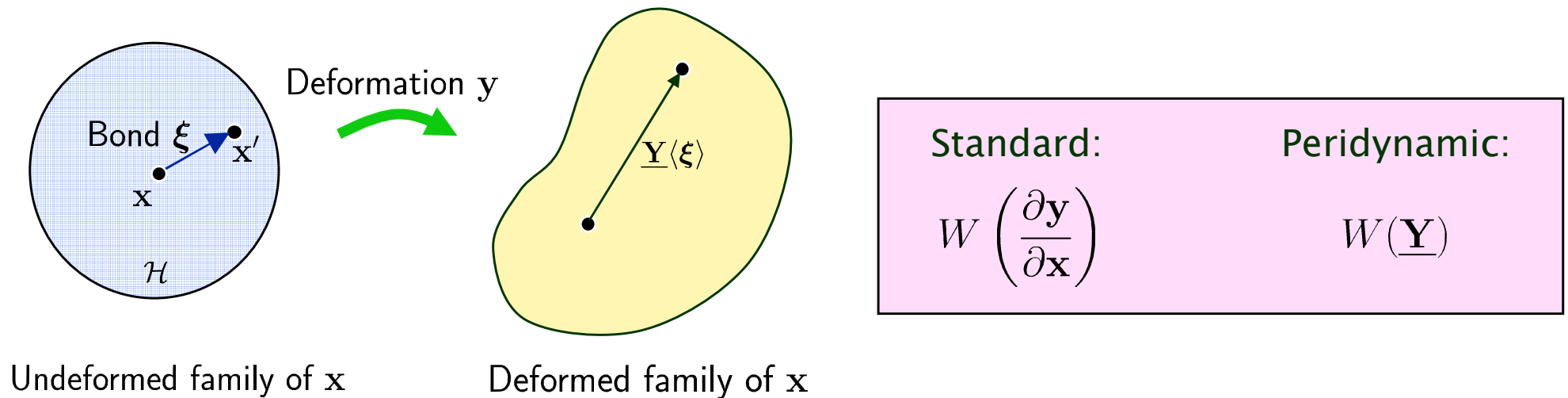
$$\begin{aligned}\Psi(\underline{\mathbf{A}} + \underline{\mathbf{a}}) &= \int f(\underline{\mathbf{A}}\langle\xi\rangle + \underline{\mathbf{a}}\langle\xi\rangle) dV_\xi \\ &= \int (f(\underline{\mathbf{A}}\langle\xi\rangle) + \text{grad } f(\underline{\mathbf{A}}\langle\xi\rangle) \cdot \underline{\mathbf{a}}\langle\xi\rangle) dV_\xi \\ &= \Psi(\underline{\mathbf{A}}) + \int \text{grad } f(\underline{\mathbf{A}}\langle\xi\rangle) \cdot \underline{\mathbf{a}}\langle\xi\rangle dV_\xi\end{aligned}$$

Therefore

$$\Psi_{\underline{\mathbf{A}}}\langle\xi\rangle = \text{grad } f(\underline{\mathbf{A}}\langle\xi\rangle).$$

Now we have the tools in place to talk about elastic materials

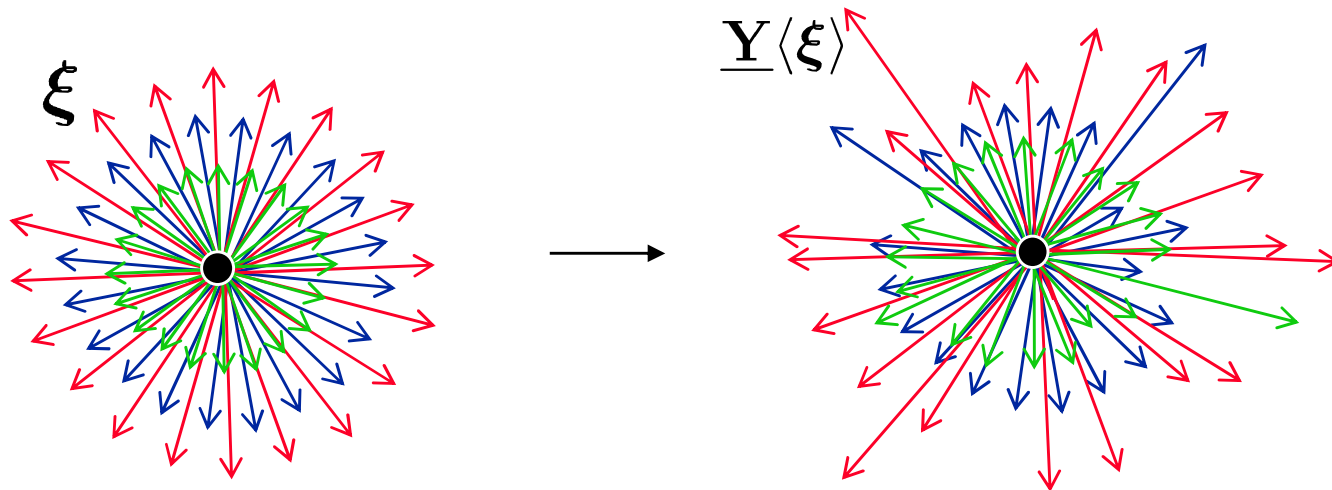
Strain energy at \mathbf{x} depends **collectively** on the deformation of the family of \mathbf{x} .



$\underline{\mathbf{Y}}$ is the *deformation state* defined by

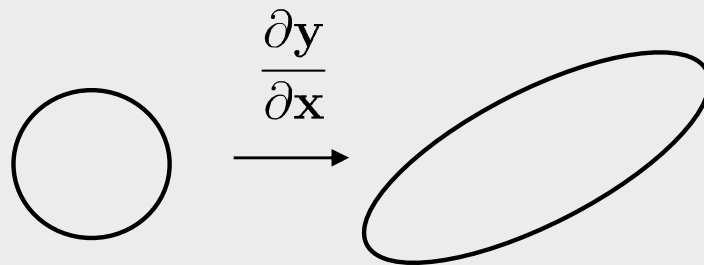
$$\underline{\mathbf{Y}}[\mathbf{x}, t]\langle \mathbf{x}' - \mathbf{x} \rangle = \mathbf{y}(\mathbf{x}', t) - \mathbf{y}(\mathbf{x}, t)$$

Deformation states can contain a lot of kinematical complexity



Undeformed bonds connected to x

Deformed bonds connected to x



Compare this with standard theory in which small spheres are mapped into ellipsoids

Force state is the work conjugate to the deformation state

- Suppose we perturb the deformed bond $\underline{\mathbf{Y}}\langle\xi\rangle$ by a virtual displacement ϵ . The resulting change in $W(\underline{\mathbf{x}})$ is

$$\Delta W = \underline{\mathbf{T}}\langle\xi\rangle \cdot \epsilon$$

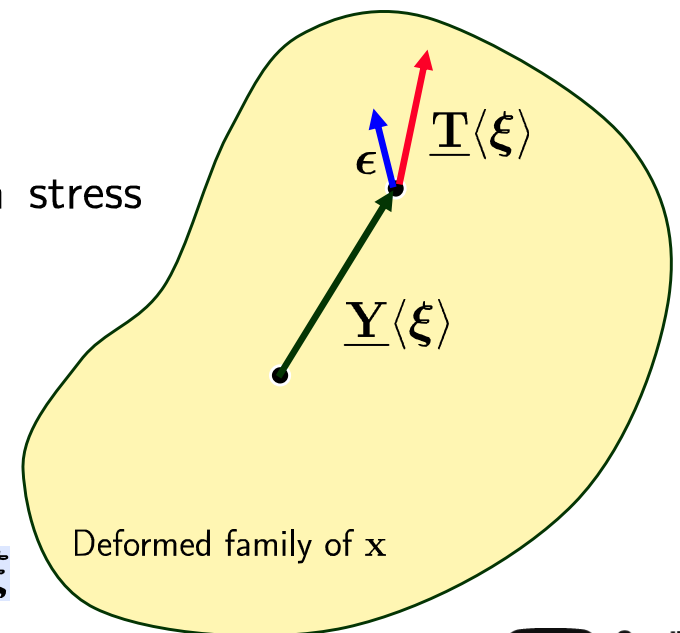
where $\underline{\mathbf{T}}\langle\xi\rangle$ is a vector.

- The “force state” $\underline{\mathbf{T}}$ is the work conjugate to $\underline{\mathbf{Y}}$:

$$\dot{W} = \underline{\mathbf{T}} \bullet \dot{\underline{\mathbf{Y}}} = \int_{\mathcal{H}} \underline{\mathbf{T}}\langle\xi\rangle \cdot \dot{\underline{\mathbf{Y}}}\langle\xi\rangle dV_{\xi}$$

- $\underline{\mathbf{T}}$ is the Frechet derivative of $W(\underline{\mathbf{Y}})$ – analogous to a stress tensor.

Displace just one bond ξ





Potential energy and its first variation

- Total potential energy in \mathcal{B} :

$$\Phi = \int_{\mathcal{B}} (W(\underline{\mathbf{Y}}[\mathbf{x}]) - \mathbf{b}(\mathbf{x}) \cdot \mathbf{y}(\mathbf{x})) dV_{\mathbf{x}}$$

- Take first variation.

$$\begin{aligned} \delta\Phi &= \int_{\mathcal{B}} (W_{\underline{\mathbf{Y}}}[\mathbf{x}] \bullet \delta\underline{\mathbf{Y}}[\mathbf{x}] - \mathbf{b}(\mathbf{x}) \cdot \delta\mathbf{y}(\mathbf{x})) dV_{\mathbf{x}} \\ &= \int_{\mathcal{B}} \left[\int_{\mathcal{B}} W_{\underline{\mathbf{Y}}}[\mathbf{x}] \langle \mathbf{x}' - \mathbf{x} \rangle \cdot (\delta\mathbf{y}(\mathbf{x}') - \delta\mathbf{y}(\mathbf{x})) dV_{\mathbf{x}'} - \mathbf{b}(\mathbf{x}) \cdot \delta\mathbf{y}(\mathbf{x}) \right] dV_{\mathbf{x}} \\ &= \int_{\mathcal{B}} \left[\int_{\mathcal{B}} (W_{\underline{\mathbf{Y}}}[\mathbf{x}'] \langle \mathbf{x} - \mathbf{x}' \rangle - W_{\underline{\mathbf{Y}}}[\mathbf{x}] \langle \mathbf{x}' - \mathbf{x} \rangle) dV_{\mathbf{x}'} - \mathbf{b}(\mathbf{x}) \right] \cdot \delta\mathbf{y}(\mathbf{x}) dV_{\mathbf{x}}. \end{aligned}$$

- Require $\delta\Phi = 0$ for all variations $\delta\mathbf{y}$. Euler-Lagrange equation is

$$\int_{\mathcal{H}} (W_{\underline{\mathbf{Y}}}[\mathbf{x}'] \langle \mathbf{x} - \mathbf{x}' \rangle - W_{\underline{\mathbf{Y}}}[\mathbf{x}] \langle \mathbf{x}' - \mathbf{x} \rangle) dV_{\mathbf{x}'} - \mathbf{b}(\mathbf{x}) = \mathbf{0}$$

for all $\mathbf{x} \in \mathcal{B}$.



Equilibrium equation

- Define the *force state* by

$$\underline{\mathbf{T}} = W_{\underline{\mathbf{Y}}}.$$

- Just showed that stationary potential energy implies the following *equilibrium equation*

$$\int_{\mathcal{H}} \left(\underline{\mathbf{T}}[\mathbf{x}] \langle \mathbf{x}' - \mathbf{x} \rangle - \underline{\mathbf{T}}[\mathbf{x}'] \langle \mathbf{x} - \mathbf{x}' \rangle \right) dV_{\mathbf{x}'} + \mathbf{b}(\mathbf{x}) = \mathbf{0}$$

for all $\mathbf{x} \in \mathcal{B}$.

Bond force

- Equilibrium equation is

$$\int_{\mathcal{H}} \left(\underline{\mathbf{T}}[\mathbf{x}] \langle \mathbf{x}' - \mathbf{x} \rangle - \underline{\mathbf{T}}[\mathbf{x}'] \langle \mathbf{x} - \mathbf{x}' \rangle \right) dV_{\mathbf{x}'} + \mathbf{b}(\mathbf{x}) = \mathbf{0}.$$

- Write this as:

$$\int_{\mathcal{H}} \mathbf{f}(\mathbf{x}', \mathbf{x}) dV_{\mathbf{x}'} + \mathbf{b}(\mathbf{x}) = \mathbf{0}.$$

- where the *bond force* is defined by

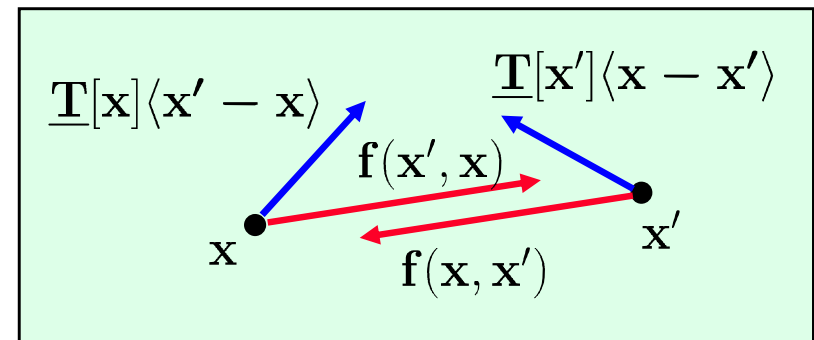
$$\mathbf{f}(\mathbf{x}', \mathbf{x}) = \underline{\mathbf{T}}[\mathbf{x}] \langle \mathbf{x}' - \mathbf{x} \rangle - \underline{\mathbf{T}}[\mathbf{x}'] \langle \mathbf{x} - \mathbf{x}' \rangle$$

- Bond force is antisymmetric:

$$\mathbf{f}(\mathbf{x}, \mathbf{x}') = -\mathbf{f}(\mathbf{x}', \mathbf{x})$$

- In general the vector $\mathbf{f}(\mathbf{x}', \mathbf{x})$ is not parallel to the deformed bond $\underline{\mathbf{Y}} \langle \mathbf{x}' - \mathbf{x} \rangle$.

- \mathbf{f} has dimensions of force/volume².



Principle of virtual work

- Equilibrium equation is

$$\int_{\mathcal{H}} \left(\underline{\mathbf{T}}[\mathbf{x}] \langle \mathbf{x}' - \mathbf{x} \rangle - \underline{\mathbf{T}}[\mathbf{x}'] \langle \mathbf{x} - \mathbf{x}' \rangle \right) dV_{\mathbf{x}'} + \mathbf{b}(\mathbf{x}) = \mathbf{0}.$$

- Multiply by a virtual displacement field \mathbf{w} and integrate:

$$\int_{\mathcal{B}} \int_{\mathcal{B}} \left(\underline{\mathbf{T}}[\mathbf{x}] \langle \mathbf{x}' - \mathbf{x} \rangle - \underline{\mathbf{T}}[\mathbf{x}'] \langle \mathbf{x} - \mathbf{x}' \rangle \right) \cdot \mathbf{w}(\mathbf{x}) dV_{\mathbf{x}'} dV_{\mathbf{x}} + \int_{\mathcal{B}} \mathbf{b}(\mathbf{x}) \cdot \mathbf{w}(\mathbf{x}) dV_{\mathbf{x}} = 0$$

$$\int_{\mathcal{B}} \int_{\mathcal{B}} \underline{\mathbf{T}}[\mathbf{x}] \langle \mathbf{x}' - \mathbf{x} \rangle \cdot (\mathbf{w}(\mathbf{x}) - \mathbf{w}(\mathbf{x}')) dV_{\mathbf{x}'} dV_{\mathbf{x}} + \int_{\mathcal{B}} \mathbf{b}(\mathbf{x}) \cdot \mathbf{w}(\mathbf{x}) dV_{\mathbf{x}} = 0$$

- If we define $\underline{\mathbf{W}}$ to be the deformation state associated with \mathbf{w}

$$\underline{\mathbf{W}}[\mathbf{x}] \langle \mathbf{x}' - \mathbf{x} \rangle = \mathbf{w}(\mathbf{x}') - \mathbf{w}(\mathbf{x})$$

then the PVW is

$$\int_{\mathcal{B}} \underline{\mathbf{T}}[\mathbf{x}] \bullet \underline{\mathbf{W}}[\mathbf{x}] dV_{\mathbf{x}} - \int_{\mathcal{B}} \mathbf{b}(\mathbf{x}) \cdot \mathbf{w}(\mathbf{x}) dV_{\mathbf{x}} = 0.$$

Compare classical PVW

$$\int (\sigma \cdot \nabla \mathbf{w} - \mathbf{b} \cdot \mathbf{w}) dV = 0$$



Peridynamic equation of motion

- Equilibrium equation:

$$\int_{\mathcal{H}} \mathbf{f}(\mathbf{x}', \mathbf{x}) dV_{\mathbf{x}'} + \mathbf{b}(\mathbf{x}) = \mathbf{0}.$$

- where

$$\mathbf{f}(\mathbf{x}', \mathbf{x}) = \underline{\mathbf{T}}[\mathbf{x}] \langle \mathbf{x}' - \mathbf{x} \rangle - \underline{\mathbf{T}}[\mathbf{x}'] \langle \mathbf{x} - \mathbf{x}' \rangle$$

- Now use d'Alembert's principle to get the equation of motion:

$$\rho(\mathbf{x}) \ddot{\mathbf{y}}(\mathbf{x}, t) = \int_{\mathcal{H}} \mathbf{f}(\mathbf{x}', \mathbf{x}, t) dV_{\mathbf{x}'} + \mathbf{b}(\mathbf{x}, t)$$

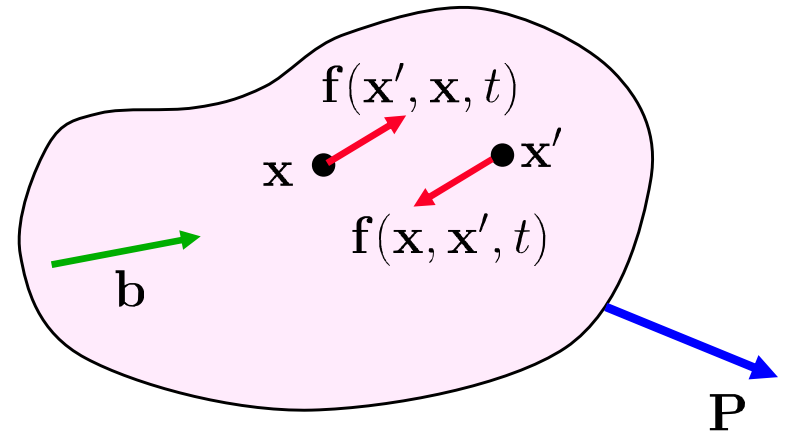
Balance of linear momentum

- Total linear momentum in the body:

$$\mathbf{P} = \int_{\mathcal{B}} \rho \dot{\mathbf{y}} dV_{\mathbf{x}}.$$

- Then

$$\begin{aligned} \dot{\mathbf{P}} &= \int_{\mathcal{B}} \rho \ddot{\mathbf{y}} dV_{\mathbf{x}} \\ &= \int_{\mathcal{B}} \left[\int_{\mathcal{B}} \mathbf{f}(\mathbf{x}', \mathbf{x}, t) dV'_{\mathbf{x}} + \mathbf{b} \right] dV_{\mathbf{x}} \end{aligned}$$



From equation of motion

- Recall $\mathbf{f}(\mathbf{x}', \mathbf{x}, t) = -\mathbf{f}(\mathbf{x}, \mathbf{x}', t)$, therefore

$$\dot{\mathbf{P}} = \int_{\mathcal{B}} \mathbf{b} dV_{\mathbf{x}}$$

- Rate of change of total momentum = total applied force.



Constitutive modeling

- A constitutive model relates the force state at a point \mathbf{x} to the deformation state and any other variables:

$$\underline{\mathbf{T}} = \hat{\underline{\mathbf{T}}}(\underline{\mathbf{Y}}, \dot{\underline{\mathbf{Y}}}, \theta, \mathbf{x}, t, \dots)$$

Diagram illustrating the variables in the constitutive model equation:

- Rate dependence (points to $\dot{\underline{\mathbf{Y}}}$)
- Temperature (points to θ)
- Heterogeneity (points to \mathbf{x})
- Explicit time dependence (e.g., ageing) (points to t)

- Simple material:

$$\underline{\mathbf{T}} = \hat{\underline{\mathbf{T}}}(\underline{\mathbf{Y}}, \mathbf{x})$$



Angular momentum balance

- Define the total angular momentum in the body by

$$\mathbf{A} = \int_{\mathcal{B}} \mathbf{y} \times \rho \dot{\mathbf{y}} dV_{\mathbf{x}}.$$

which says there are no “hidden” dofs that have angular momentum.

- Shorten the notation:

$$\mathbf{t} = \underline{\mathbf{T}}[\mathbf{x}, t] \langle \mathbf{x}' - \mathbf{x} \rangle, \quad \mathbf{t}' = \underline{\mathbf{T}}[\mathbf{x}', t] \langle \mathbf{x} - \mathbf{x}' \rangle.$$

- Then

$$\begin{aligned} \dot{\mathbf{A}} &= \int_{\mathcal{B}} \mathbf{y} \times \rho \ddot{\mathbf{y}} dV_{\mathbf{x}} \\ &= \int_{\mathcal{B}} \mathbf{y} \times \left[\int_{\mathcal{B}} (\mathbf{t} - \mathbf{t}') dV_{\mathbf{x}'} + \mathbf{b} \right] dV_{\mathbf{x}} \end{aligned}$$

From equation of motion



Angular momentum balance: Nonpolar materials

$$\begin{aligned}\dot{\mathbf{A}} &= \int_{\mathcal{B}} \int_{\mathcal{B}} \mathbf{y} \times (\mathbf{t} - \mathbf{t}') dV_{\mathbf{x}'} dV_{\mathbf{x}} + \int_{\mathcal{B}} \mathbf{y} \times \mathbf{b} dV_{\mathbf{x}} \\ &= \int_{\mathcal{B}} \int_{\mathcal{B}} (\mathbf{y} - \mathbf{y}') \times \mathbf{t} dV_{\mathbf{x}'} dV_{\mathbf{x}} + \int_{\mathcal{B}} \mathbf{y} \times \mathbf{b} dV_{\mathbf{x}} \\ &= - \int_{\mathcal{B}} \int_{\mathcal{H}} \underline{\mathbf{Y}}\langle \boldsymbol{\xi} \rangle \times \underline{\mathbf{T}}\langle \boldsymbol{\xi} \rangle dV_{\boldsymbol{\xi}} dV_{\mathbf{x}} + \int_{\mathcal{B}} \mathbf{y} \times \mathbf{b} dV_{\mathbf{x}}\end{aligned}$$

Suppose the constitutive model is *nonpolar*:

$$\int_{\mathcal{H}} \underline{\mathbf{Y}}\langle \boldsymbol{\xi} \rangle \times \underline{\mathbf{T}}\langle \boldsymbol{\xi} \rangle dV_{\boldsymbol{\xi}} = \mathbf{0}$$

for all $\underline{\mathbf{Y}}$. Then

$$\dot{\mathbf{A}} = \int_{\mathcal{B}} \mathbf{y} \times \mathbf{b} dV_{\mathbf{x}}$$

which says there are no “hidden” moments.

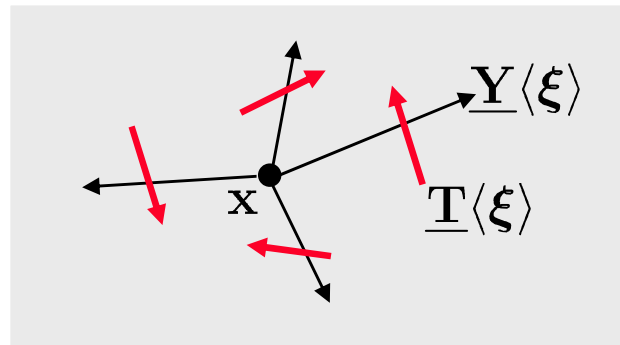
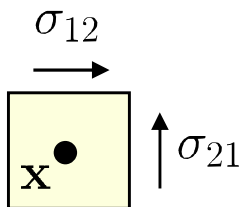
Nonpolar materials

- Nonpolarity:

$$\int_{\mathcal{H}} \underline{\mathbf{Y}}\langle \boldsymbol{\xi} \rangle \times \hat{\mathbf{T}}(\underline{\mathbf{Y}})\langle \boldsymbol{\xi} \rangle dV_{\boldsymbol{\xi}} = \mathbf{0} \quad \forall \underline{\mathbf{Y}}$$

implies the global balance of angular momentum.

- Converse can be proved too (global balance of angular momentum implies material is nonpolar).
- We will adopt nonpolarity as a constitutive restriction.
- “No net moment on a point due to its own force state.”



Ordinary and nonordinary

- Any force state can be decomposed into parts that are parallel and orthogonal to the deformed bonds:

$$\underline{\mathbf{T}} = \underline{\mathbf{T}}_{\parallel} + \underline{\mathbf{T}}_{\perp}$$

where

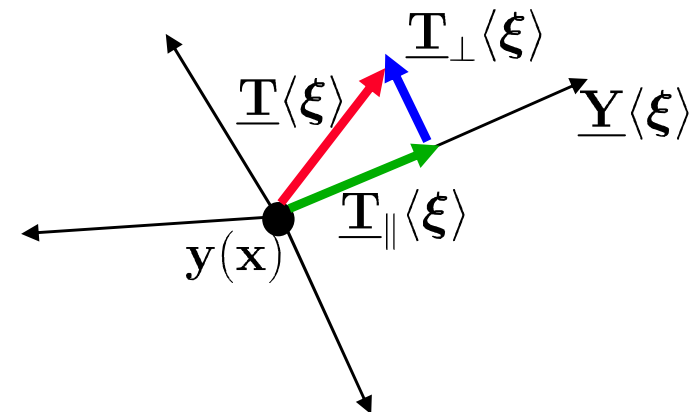
$$\underline{\mathbf{T}}_{\parallel} \langle \xi \rangle = (\underline{\mathbf{T}} \langle \xi \rangle \cdot \underline{\mathbf{M}} \langle \xi \rangle) \underline{\mathbf{M}} \langle \xi \rangle$$


$$\underline{\mathbf{M}} \langle \xi \rangle = \frac{\underline{\mathbf{Y}} \langle \xi \rangle}{|\underline{\mathbf{Y}} \langle \xi \rangle|}.$$

- If

$$\underline{\mathbf{T}}_{\perp} = \underline{\mathbf{0}} \quad \forall \underline{\mathbf{Y}}$$

then the material is *ordinary*, otherwise *nonordinary*.





Elastic materials: objectivity implies nonpolarity

- Objectivity: for any proper orthogonal tensor \mathbf{Q} ,

$$W(\mathbf{Q}\underline{\mathbf{Y}}) = W(\underline{\mathbf{Y}}).$$

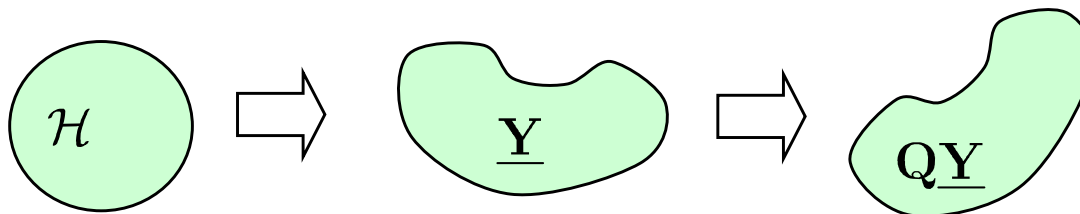
i.e., energy doesn't change if you rigidly rotate the family after deforming it.

- Can show that any objective, elastic material is nonpolar:

$$\int_{\mathcal{H}} \underline{\mathbf{Y}}(\underline{\boldsymbol{\xi}}) \times W_{\underline{\mathbf{Y}}}(\underline{\boldsymbol{\xi}}) dV_{\boldsymbol{\xi}} = \mathbf{0}.$$

Details: see Silling, "Linearized theory of peridynamic states," J. Elast. (2010).

- Result is important because usually objectivity is much easier to determine than nonpolarity directly.



Energy balance

- Recall that for an elastic material, since $\underline{\mathbf{T}} = W_{\underline{\mathbf{Y}}}$,

$$W(\underline{\mathbf{Y}} + \delta \underline{\mathbf{Y}}) = W(\underline{\mathbf{Y}}) + \underline{\mathbf{T}} \bullet \delta \underline{\mathbf{Y}} + o(||\delta \underline{\mathbf{Y}}||)$$

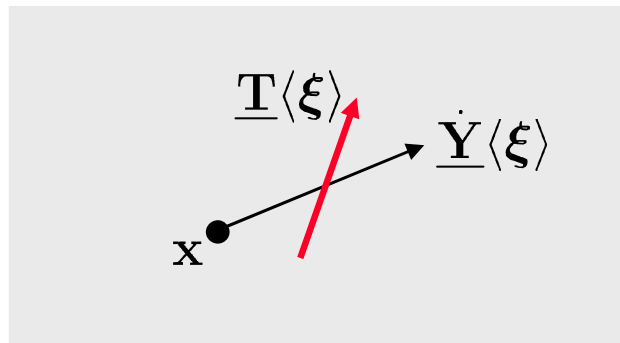
therefore

$$\dot{W} = \underline{\mathbf{T}} \bullet \dot{\underline{\mathbf{Y}}}$$

i.e.,

$$\dot{W}(\mathbf{x}, t) = \int_{\mathcal{H}} \underline{\mathbf{T}}[\mathbf{x}, t] \langle \mathbf{x}' - \mathbf{x} \rangle \cdot (\dot{\mathbf{y}}(\mathbf{x}', t) - \dot{\mathbf{y}}(\mathbf{x}, t)) dV_{\mathbf{x}'}$$

Compare stress power: $\dot{W} = \boldsymbol{\sigma} \cdot \dot{\mathbf{F}}$.





Energy balance, ctd.

- For more general materials, the first law of thermodynamics is

$$\dot{\varepsilon} = \underline{\mathbf{T}} \bullet \underline{\dot{\mathbf{Y}}} + h + r$$

where ε =internal energy density, h =net heat transport rate to \mathbf{x} per unit volume, r = heat source rate.

- This applies to any heat transport law, e.g.

$$h = K \nabla^2 \theta \quad \text{Fourier's law, local}$$

or

$$h = \int_{\mathcal{B}} K(\mathbf{x}' - \mathbf{x})(\theta(\mathbf{x}', t) - \theta(\mathbf{x}, t)) dV_{\mathbf{x}'} \quad \text{nonlocal}$$



Free energy and 2nd law of thermodynamics

(joint work with Rich Lehoucq, thanks also to Eliot Fried)

- Since we're now dealing with temperature, have to include it in the internal energy:

$$\varepsilon(\underline{\mathbf{Y}}, \theta).$$

- Now try to find $\underline{\mathbf{T}}$ from ε . Define the *free energy* by

$$\psi = \varepsilon - \theta \eta$$

where η =entropy.

- Thus

$$\dot{\psi} = \dot{\varepsilon} - \dot{\theta} \eta - \theta \dot{\eta}.$$

- Hence from 1st law

$$\dot{\psi} = \underline{\mathbf{T}} \bullet \underline{\dot{\mathbf{Y}}} + h + r - \dot{\theta} \eta - \theta \dot{\eta}.$$

- Second law (Clausius inequality):

$$\theta \dot{\eta} \geq h + r.$$



Free energy and the force state

- From last two equations,

$$\underline{\mathbf{T}} \bullet \underline{\dot{\mathbf{Y}}} - \dot{\theta} \eta - \dot{\psi} \geq 0.$$

- Assume $\psi = \psi(\underline{\mathbf{Y}}, \theta)$. Therefore

$$\underline{\mathbf{T}} \bullet \underline{\dot{\mathbf{Y}}} - \dot{\theta} \eta - (\psi_{\underline{\mathbf{Y}}} \bullet \underline{\dot{\mathbf{Y}}} + \psi_{\theta} \dot{\theta}) \geq 0.$$

- Group terms:

$$(\underline{\mathbf{T}} - \psi_{\underline{\mathbf{Y}}}) \bullet \underline{\dot{\mathbf{Y}}} - (\eta + \psi_{\theta}) \dot{\theta} \geq 0.$$

- Since $\underline{\mathbf{Y}}$ and θ can (in principle) be varied independently, conclude (from Coleman-Noll argument) that

$$\underline{\mathbf{T}} = \psi_{\underline{\mathbf{Y}}} \quad \text{and} \quad \eta = -\psi_{\theta}.$$

- Special case: if ψ is independent of θ , get an elastic material with $W = \psi$.



Rate dependent materials

- If we allow rate dependence in the model,

$$\psi = \psi(\underline{\mathbf{Y}}, \underline{\dot{\mathbf{Y}}}, \theta)$$

can show that the force state can be decomposed into *equilibrium* and *dissipative* parts:

$$\underline{\mathbf{T}} = \underline{\mathbf{T}}^e(\underline{\mathbf{Y}}, \theta) + \underline{\mathbf{T}}^d(\underline{\mathbf{Y}}, \underline{\dot{\mathbf{Y}}}, \theta)$$

where

$$\underline{\mathbf{T}}^e = \psi_{\underline{\mathbf{Y}}} \quad \text{and} \quad \underline{\mathbf{T}}^d \bullet \underline{\dot{\mathbf{Y}}} \geq 0.$$

↑
Energy dissipation

Material modeling: Bond-based materials

- The simplest assumption is that all the bonds are independent.

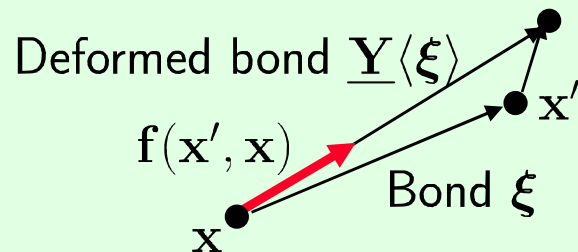
$$W(\underline{\mathbf{Y}}) = \int_{\mathcal{H}} \psi(\underline{e}(\underline{\xi}), \underline{\xi}) dV_{\xi}, \quad \underline{e}(\underline{\xi}) = |\underline{\mathbf{Y}}(\underline{\xi})| - |\underline{\xi}|$$

$$\underline{\mathbf{T}}(\underline{\xi}) = \psi'(\underline{e}(\underline{\xi}), \underline{\xi}) \mathbf{M}, \quad \mathbf{M} = \frac{\underline{\mathbf{Y}}(\underline{\xi})}{|\underline{\mathbf{Y}}(\underline{\xi})|}$$

- Equation of motion simplifies to

$$\rho \ddot{\mathbf{y}}(\mathbf{x}, t) = \int_{\mathcal{H}} \mathbf{f}(\mathbf{x}', \mathbf{x}) dV_{\mathbf{x}} + \mathbf{b}(\mathbf{x}, t),$$

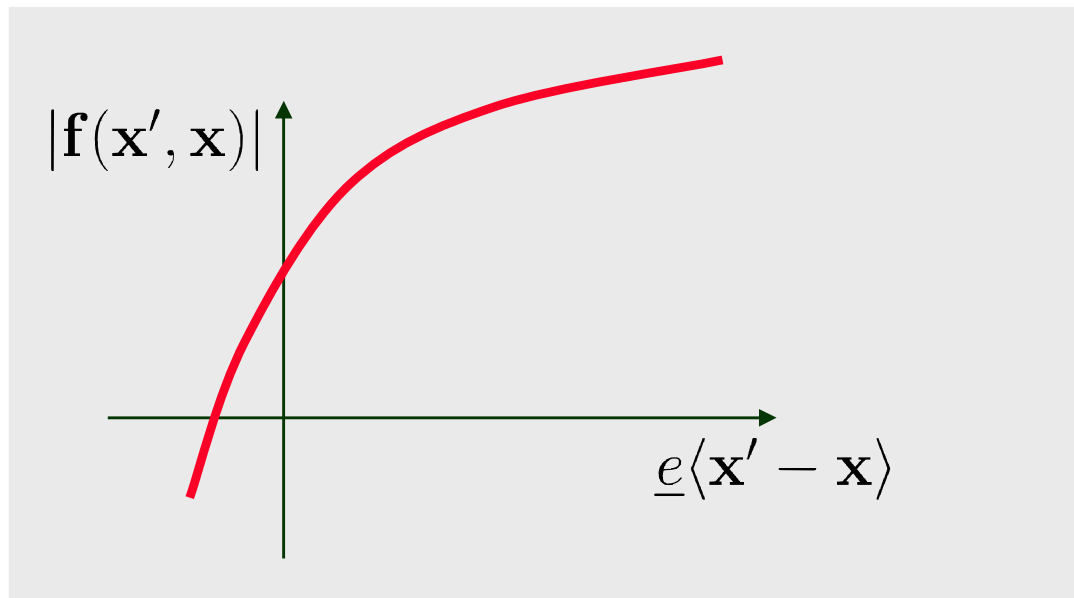
$$\mathbf{f}(\mathbf{x}', \mathbf{x}) = (\psi'(\underline{e}(\underline{\xi}), \underline{\xi}) + \psi'(\underline{e}(\underline{\xi}), -\underline{\xi})) \mathbf{M}$$



$$\text{Bond extension} = \underline{e}(\underline{\xi}) = |\underline{\mathbf{Y}}(\underline{\xi})| - |\underline{\xi}|$$

Material modeling: Bond-based materials, ctd.

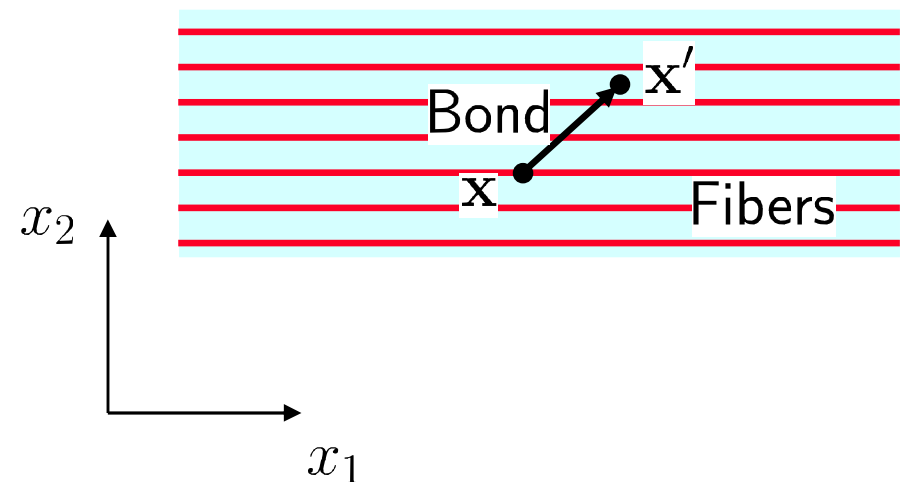
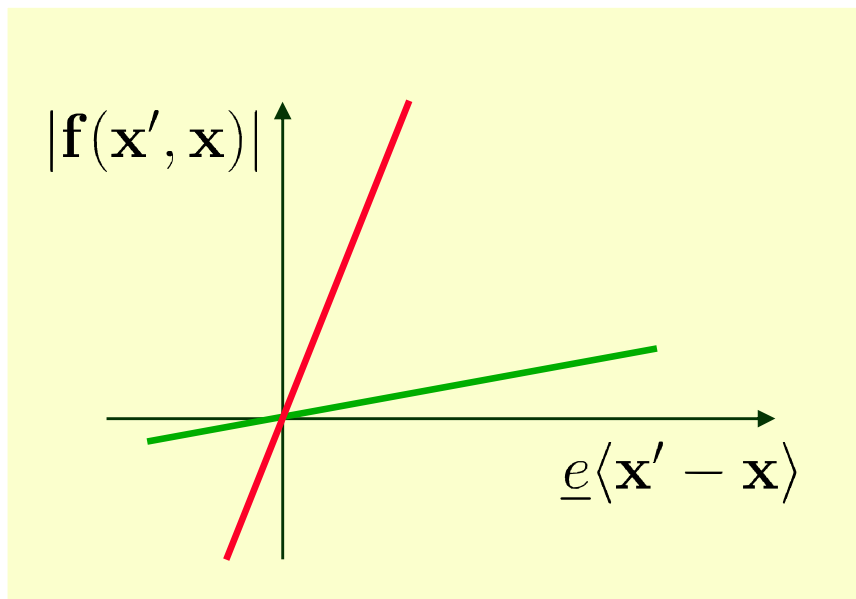
- The body is a network of independent, nonlinear springs.
- Material response is described by a graph of bond force vs. bond extension.
- If the material is isotropic, the Poisson ratio = $1/4$ (!).



Material modeling: Bond-based materials, ctd.

- Special case: fiber reinforced composite lamina.
- Bonds in the fiber direction are stiffer than the others.

$$\mathbf{f}(\mathbf{x}', \mathbf{x}) = (c_1 + c_2 \Delta(x'_2 - x_2)) \underline{e} \langle \mathbf{x}' - \mathbf{x} \rangle \mathbf{M}$$





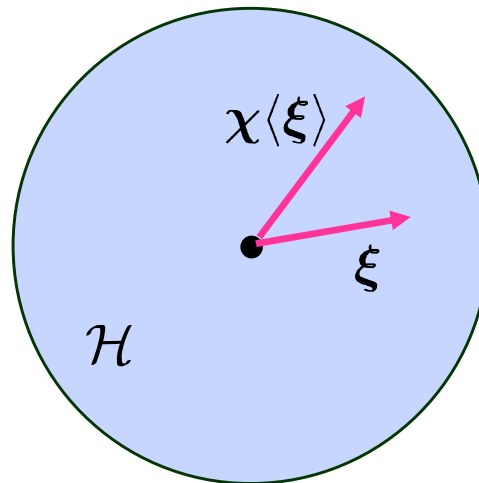
Material modeling: Bond-pair materials

- Suppose every bond ξ has a friend $\eta = \chi(\xi)$. The material responds to the deformation of pairs of bonds.

$$W(\underline{\mathbf{Y}}) = \int_{\mathcal{H}} \psi(\underline{\mathbf{Y}}\langle\xi\rangle, \underline{\mathbf{Y}}\langle\eta\rangle, \xi, \eta) dV_{\xi}$$

where ψ is a function of four vectors:

$$\psi(\mathbf{p}, \mathbf{q}, \xi, \eta).$$



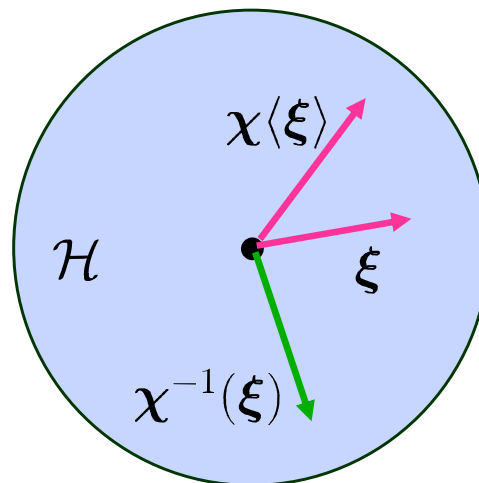
Material modeling: Bond-pair materials, ctd.

- Fréchet derivative yields

$$\underline{\mathbf{T}}\langle \xi \rangle = \psi_p(\underline{\mathbf{Y}}\langle \xi \rangle, \underline{\mathbf{Y}}\langle \chi(\xi) \rangle, \xi, \chi(\xi)) + \psi_q(\underline{\mathbf{Y}}\langle \chi^{-1}(\xi) \rangle, \underline{\mathbf{Y}}\langle \xi \rangle, \chi^{-1}(\xi), \xi) J^{-1}$$

where

$$J = |\det \text{grad } \chi|$$



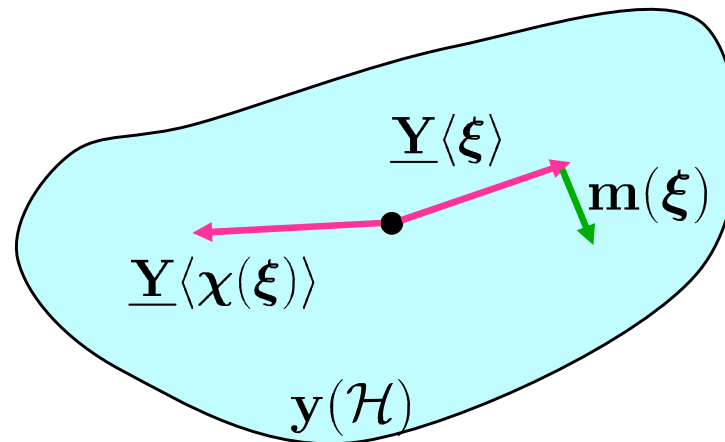
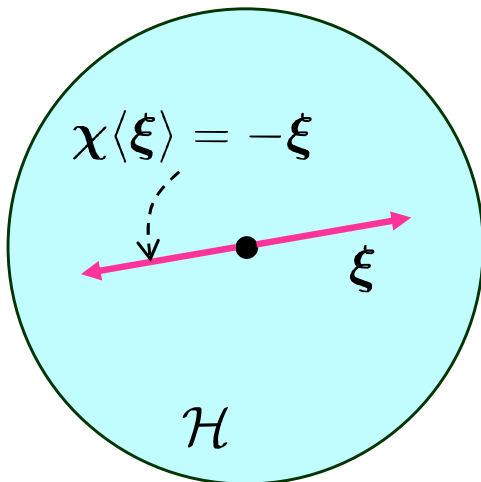
Material modeling: Bond-pair materials, ctd.

- Specific case:

$$\chi(\xi) = -\xi$$

$$\psi(\underline{\mathbf{Y}}\langle\xi\rangle, \underline{\mathbf{Y}}\langle\eta\rangle, \xi, \eta) = \frac{c}{4}(\theta - \pi)^2, \quad \theta = \cos^{-1} \frac{\underline{\mathbf{Y}}\langle\xi\rangle \cdot \underline{\mathbf{Y}}\langle\eta\rangle}{|\underline{\mathbf{Y}}\langle\xi\rangle| |\underline{\mathbf{Y}}\langle\eta\rangle|}$$

$$\underline{\mathbf{T}}\langle\xi\rangle = \frac{c(\pi - \theta)}{|\underline{\mathbf{Y}}\langle\xi\rangle|} \mathbf{m}(\xi)$$

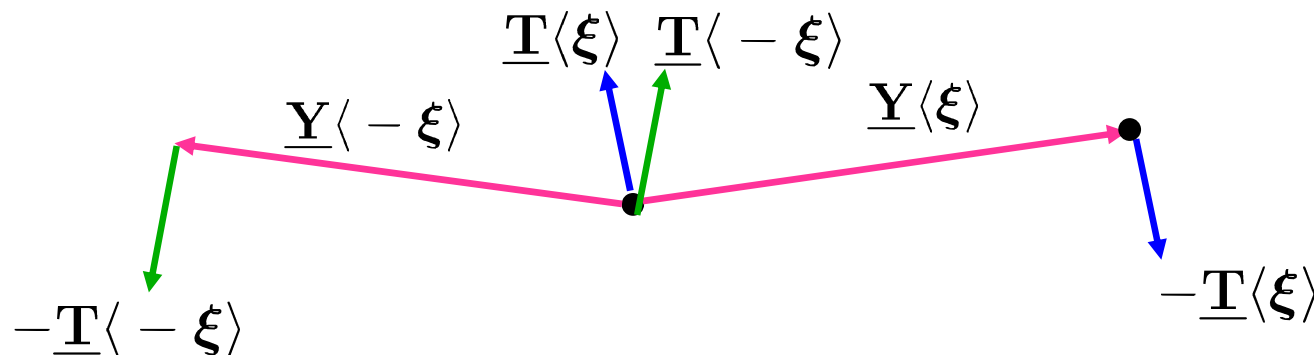


$\mathbf{m}(\xi)$ = unit vector \perp to $\underline{\mathbf{Y}}\langle\xi\rangle$

Material modeling: Bond-pair materials, ctd.

Fascinating facts:

- This material does not respond at all to homogeneous deformation.
- It provides a consistent way to model bending of a one-dimensional beam.
- The standard model for a beam involves introducing a different theory from the continuum theory.



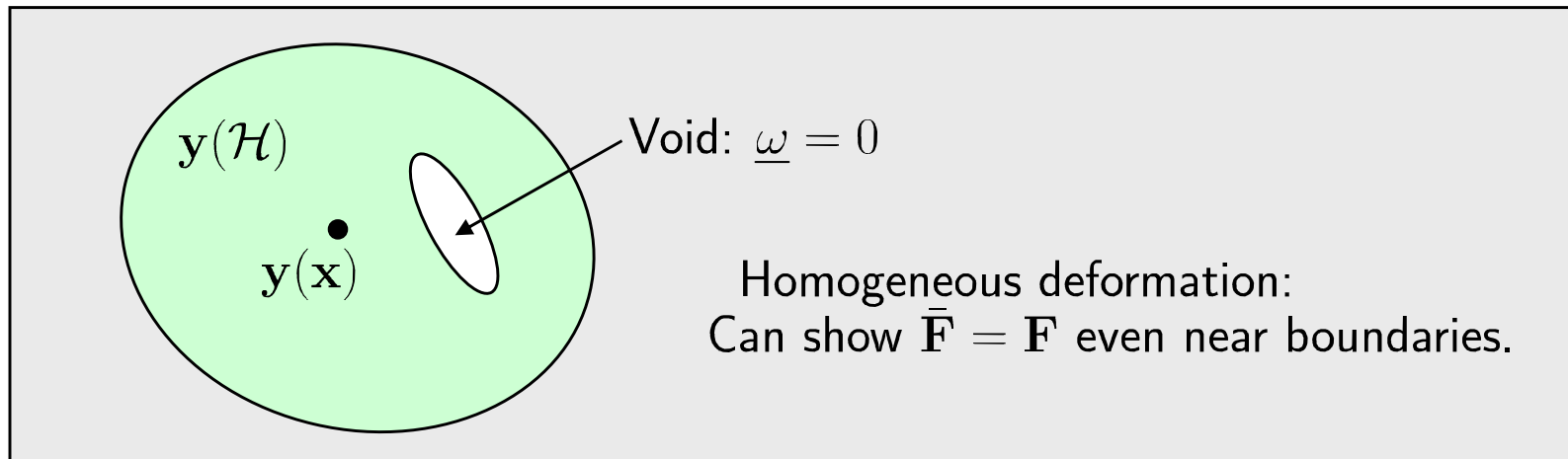
Using a classical stress–strain material model in peridynamics

- Suppose we are given a model for the Piola stress: $\sigma(\mathbf{F})$, where $\mathbf{F} = \nabla \mathbf{y}$.
- Want to use this somehow to get a force state. Define an approximate \mathbf{F} by

$$\bar{\mathbf{F}} = \left[\int_{\mathcal{H}} \underline{\omega}(\boldsymbol{\xi}) \underline{\mathbf{Y}}(\boldsymbol{\xi}) \otimes \boldsymbol{\xi} dV_{\boldsymbol{\xi}} \right] \mathbf{K}^{-1}$$

where $\underline{\omega}$ is a given *influence function* and \mathbf{K} is the *shape tensor*:

$$\mathbf{K} = \int_{\mathcal{H}} \underline{\omega}(\boldsymbol{\xi}) \boldsymbol{\xi} \otimes \boldsymbol{\xi} dV_{\boldsymbol{\xi}}$$



Using a classical stress–strain material model in peridynamics, ctd.

- Set

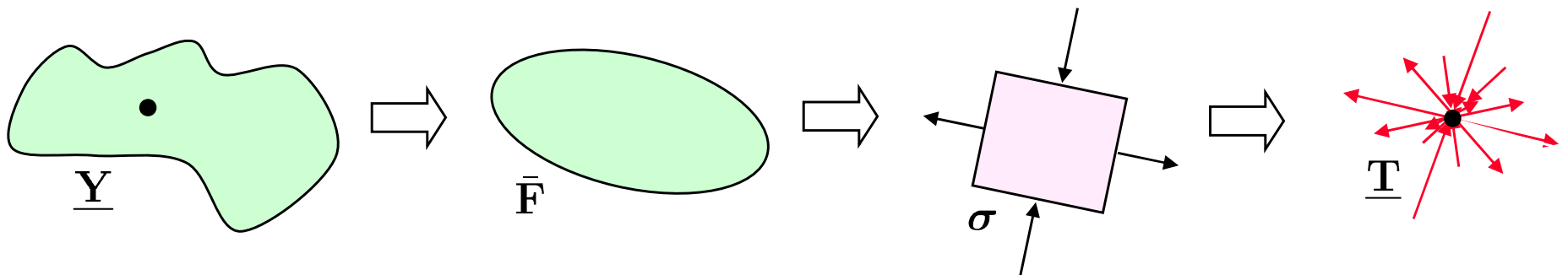
$$\underline{\mathbf{T}}\langle \underline{\boldsymbol{\xi}} \rangle = \underline{\omega}\langle \underline{\boldsymbol{\xi}} \rangle \boldsymbol{\sigma}(\bar{\mathbf{F}}) \mathbf{K}^{-1} \underline{\boldsymbol{\xi}} \quad \forall \underline{\boldsymbol{\xi}}$$

- Can show that if $\boldsymbol{\sigma} = \partial W / \partial \mathbf{F}$, then $\underline{\mathbf{T}} = W_{\underline{\mathbf{Y}}}$.
- Can also show that if the Cauchy stress tensor is symmetric, i.e.,

$$\boldsymbol{\tau}^T = \boldsymbol{\tau} \quad \text{where} \quad \boldsymbol{\tau} = \frac{\boldsymbol{\sigma}(\mathbf{F}) \mathbf{F}^T}{\det \mathbf{F}}$$

then the peridynamic material is nonpolar:

$$\int_{\mathcal{H}} \underline{\mathbf{Y}}\langle \underline{\boldsymbol{\xi}} \rangle \times \underline{\mathbf{T}}\langle \underline{\boldsymbol{\xi}} \rangle dV_{\underline{\boldsymbol{\xi}}} = \mathbf{0} \quad \forall \underline{\mathbf{Y}}.$$





Fluids

- Define a nonlocal dilatation based on the mean bond extension:

$$\vartheta = \frac{3}{m} \underline{\omega x} \bullet \underline{e}$$

where

$$\underline{x}\langle \underline{\xi} \rangle = |\underline{\xi}|, \quad m = \underline{\omega x} \bullet \underline{x}, \quad \underline{e}\langle \underline{\xi} \rangle = |\underline{Y}\langle \underline{\xi} \rangle| - \underline{x}\langle \underline{\xi} \rangle.$$

- Writing this out in detail:

$$\vartheta = \frac{3}{m} \int_{\mathcal{H}} \underline{\omega}\langle \underline{\xi} \rangle |\underline{\xi}| (|\underline{Y}\langle \underline{\xi} \rangle| - |\underline{\xi}|) dV_{\underline{\xi}}.$$

- Constitutive model: $W(\vartheta)$.

Fluids, ctd.

- Nonlocal dilatation:

$$\vartheta(\underline{\mathbf{Y}}) = \frac{3}{m} \int_{\mathcal{H}} \underline{\omega}(\underline{\xi}) |\underline{\xi}| (|\underline{\mathbf{Y}}(\underline{\xi})| - |\underline{\xi}|) dV_{\underline{\xi}}.$$

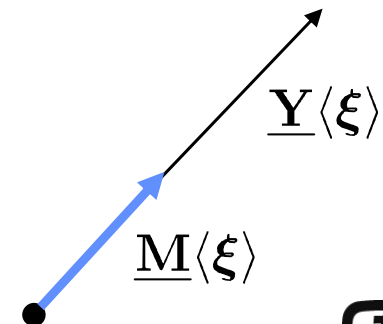
- Fréchet derivative of dilatation: observe

$$\begin{aligned} \vartheta(\underline{\mathbf{Y}} + \delta \underline{\mathbf{Y}}) &= \frac{3}{m} \int_{\mathcal{H}} \underline{\omega}(\underline{\xi}) |\underline{\xi}| (|\underline{\mathbf{Y}}(\underline{\xi}) + \delta \underline{\mathbf{Y}}(\underline{\xi})| - |\underline{\xi}|) dV_{\underline{\xi}} \\ &= \frac{3}{m} \int_{\mathcal{H}} \underline{\omega}(\underline{\xi}) |\underline{\xi}| \left(|\underline{\mathbf{Y}}(\underline{\xi})| + \frac{\underline{\mathbf{Y}}(\underline{\xi})}{|\underline{\mathbf{Y}}(\underline{\xi})|} \cdot \delta \underline{\mathbf{Y}}(\underline{\xi}) - |\underline{\xi}| \right) dV_{\underline{\xi}} \\ &= \vartheta(\underline{\mathbf{Y}}) + \frac{3}{m} \int_{\mathcal{H}} \underline{\omega}(\underline{\xi}) |\underline{\xi}| \left(\frac{\underline{\mathbf{Y}}(\underline{\xi})}{|\underline{\mathbf{Y}}(\underline{\xi})|} \cdot \delta \underline{\mathbf{Y}}(\underline{\xi}) \right) dV_{\underline{\xi}} \end{aligned}$$

hence

$$\vartheta_{\underline{\mathbf{Y}}} = \frac{3}{m} \underline{\omega x M} \quad \text{where} \quad \underline{\mathbf{M}} = \frac{\underline{\mathbf{Y}}}{|\underline{\mathbf{Y}}|}.$$

$\underline{\mathbf{M}}(\underline{\xi})$ is the deformed bond direction

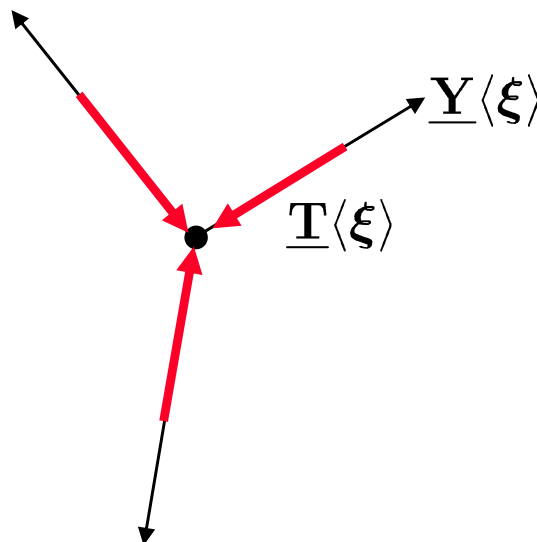


Fluids, ctd.

- Constitutive model is $W(\vartheta)$.
- Now can write down the force state: chain rule implies

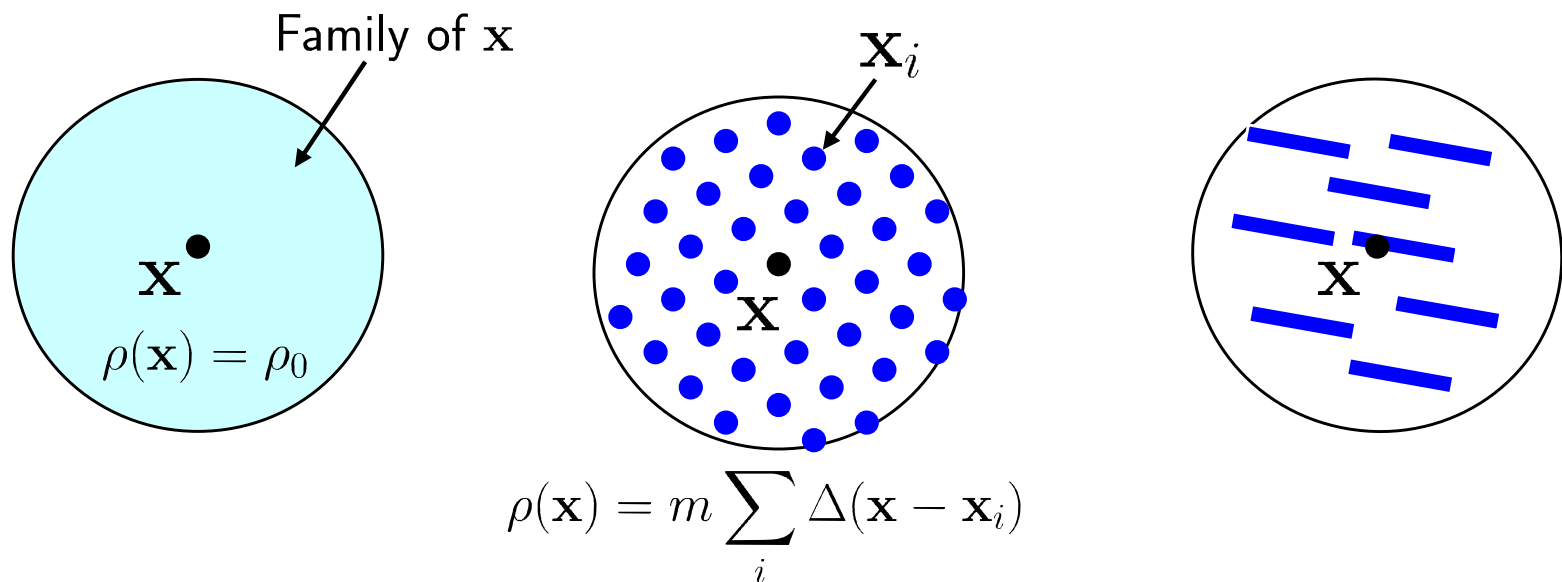
$$\hat{\underline{\mathbf{T}}}(\underline{\mathbf{Y}}) = W_{\underline{\mathbf{Y}}}(\vartheta(\underline{\mathbf{Y}})) = \frac{dW}{d\vartheta}(\vartheta(\underline{\mathbf{Y}}))\vartheta_{\underline{\mathbf{Y}}} = \frac{dW}{d\vartheta}(\vartheta(\underline{\mathbf{Y}})) \frac{3\omega x \underline{\mathbf{M}}}{m}.$$

- Nonlocal pressure $= -dW/d\vartheta$.
- Bond forces are parallel to the deformed bonds (material is ordinary).



Material modeling: Discrete particles

- The family of \mathbf{x} could be either continuous or a collection of point masses or other objects.



$\Delta = 3\text{D Dirac delta function}$



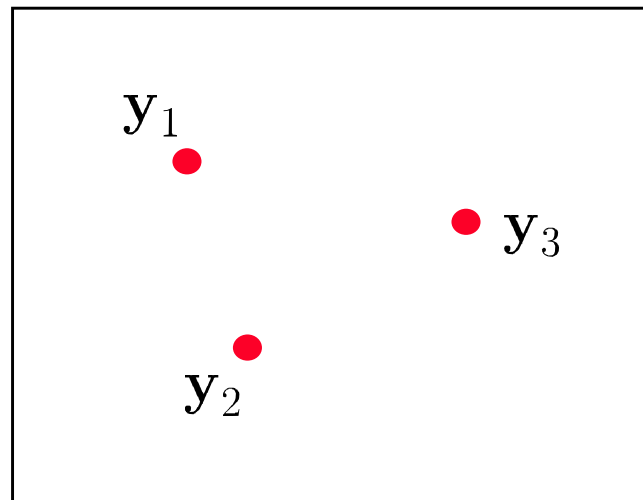
Material modeling: Discrete particles, ctd.

- Consider a set of atoms that interact through an N -body potential:

$$U(\mathbf{y}_1, \mathbf{y}_2, \dots, \mathbf{y}_N),$$

$\mathbf{y}_1, \dots, \mathbf{y}_N$ = deformed positions, $\mathbf{x}_1, \dots, \mathbf{x}_N$ = reference positions.

- This can be represented exactly as a peridynamic body.

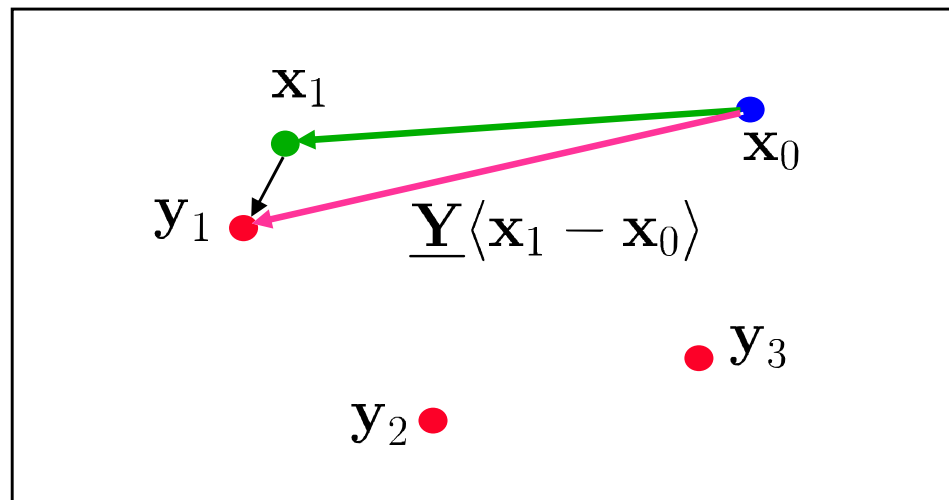


Material modeling: Discrete particles, ctd.

Define a peridynamic body by:

$$\hat{W}(\underline{\mathbf{Y}}, \mathbf{x}) = \Delta(\mathbf{x} - \mathbf{x}_0) U(\underline{\mathbf{Y}}\langle \mathbf{x}_1 - \mathbf{x}_0 \rangle, \underline{\mathbf{Y}}\langle \mathbf{x}_2 - \mathbf{x}_0 \rangle, \dots, \underline{\mathbf{Y}}\langle \mathbf{x}_N - \mathbf{x}_0 \rangle),$$

$$\rho(\mathbf{x}) = \sum_i \Delta(\mathbf{x} - \mathbf{x}_i) M_i$$



Material modeling: Discrete particles, ctd.

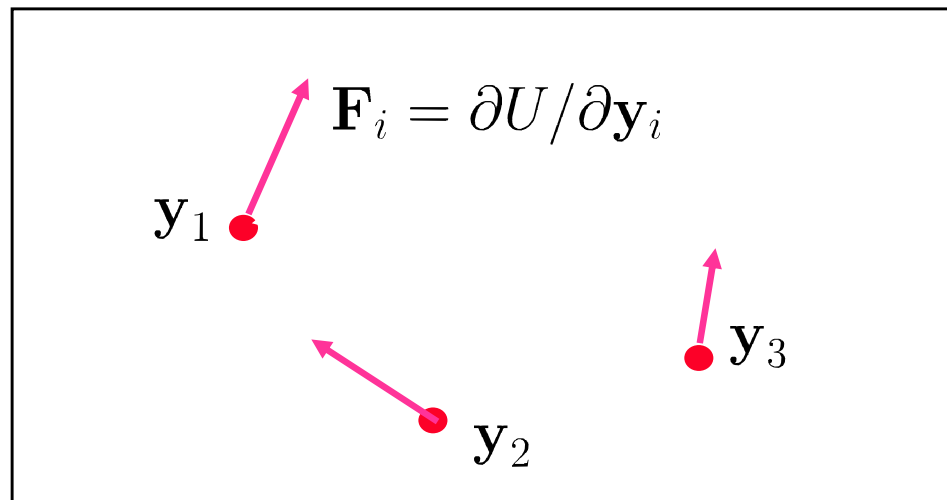
- After evaluating the Frechet derivative $\underline{\mathbf{T}}$, find

$$\underline{\mathbf{T}}[\mathbf{x}]\langle \boldsymbol{\xi} \rangle = \Delta(\mathbf{x} - \mathbf{x}_0) \sum_i \frac{\partial U}{\partial \mathbf{y}_i} \Delta(\boldsymbol{\xi} - (\mathbf{x}_i - \mathbf{x}_0))$$

- Equation of motion reduces to

$$M_i \ddot{\mathbf{y}}(\mathbf{x}_i, t) = -\frac{\partial U}{\partial \mathbf{y}_i}, \quad i = 1, \dots, N$$

- Have represented a multibody potential exactly within a continuum model.





Linearization of a material model

- Small displacement field $\underline{\mathbf{u}}$ superposed on a (possibly) large deformation \mathbf{y}^0 :

$$\hat{\underline{\mathbf{T}}}(\underline{\mathbf{Y}}^0 + \underline{\mathbf{U}}) = \hat{\underline{\mathbf{T}}}(\underline{\mathbf{Y}}^0) + \underline{\mathbb{K}} \bullet \underline{\mathbf{U}} + o(||\underline{\mathbf{U}}||)$$

where

$$\underline{\mathbf{Y}}^0 \langle \mathbf{x}' - \mathbf{x} \rangle = \mathbf{y}(\mathbf{x}') - \mathbf{y}(\mathbf{x})$$

$$\underline{\mathbf{U}} \langle \mathbf{x}' - \mathbf{x} \rangle = \mathbf{u}(\mathbf{x}') - \mathbf{u}(\mathbf{x})$$

$$\underline{\mathbb{K}} = \hat{\underline{\mathbf{T}}}_{\underline{\mathbf{Y}}}(\underline{\mathbf{Y}}^0)$$

- $\underline{\mathbb{K}} \langle \boldsymbol{\xi}, \boldsymbol{\zeta} \rangle$ is a *double state* (tensor valued function of two bonds):

$$(\underline{\mathbb{K}} \bullet \underline{\mathbf{U}}) \langle \boldsymbol{\xi} \rangle = \int_{\mathcal{H}} \underline{\mathbb{K}} \langle \boldsymbol{\xi}, \boldsymbol{\zeta} \rangle \underline{\mathbf{U}} \langle \boldsymbol{\zeta} \rangle dV_{\boldsymbol{\zeta}}$$



Linearization of an elastic material model

- If $\hat{\underline{\mathbf{T}}}$ is elastic,

$$\underline{\mathbb{K}} = \hat{\underline{\mathbf{T}}}_{\underline{\mathbf{Y}}}(\underline{\mathbf{Y}}^0) = W_{\underline{\mathbf{Y}}\underline{\mathbf{Y}}}(\underline{\mathbf{Y}}^0)$$

i.e., $\underline{\mathbb{K}}$ is the second Fréchet derivative of W .

- Can show that for a linearized elastic material,

$$\underline{\mathbb{K}}\langle \underline{\zeta}, \underline{\xi} \rangle = \underline{\mathbb{K}}^T \langle \underline{\xi}, \underline{\zeta} \rangle \quad \forall \underline{\xi}, \underline{\zeta}$$

- Converse is also true.
- $\underline{\mathbb{K}}$ is called the *modulus state*, similar to 4th order elasticity tensor.



Equation of motion for a linearized material

- If \mathbf{y}^0 is equilibrated,

$$\begin{aligned}\rho \ddot{\mathbf{u}}(\mathbf{x}) &= \int (\underline{\mathbf{T}}[\mathbf{x}] \langle \mathbf{p} - \mathbf{x} \rangle - \underline{\mathbf{T}}[\mathbf{p}] \langle \mathbf{x} - \mathbf{p} \rangle) dV_{\mathbf{p}} + \mathbf{b}(\mathbf{x}) \\ &= \int ((\underline{\mathbb{K}}[\mathbf{x}] \bullet \underline{\mathbf{U}}[\mathbf{x}]) \langle \mathbf{p} - \mathbf{x} \rangle - (\underline{\mathbb{K}}[\mathbf{p}] \bullet \underline{\mathbf{U}}[\mathbf{p}]) \langle \mathbf{x} - \mathbf{p} \rangle) dV_{\mathbf{p}} + \mathbf{b}(\mathbf{x}) \\ &= \int \int (\underline{\mathbb{K}}[\mathbf{x}] \langle \mathbf{p} - \mathbf{x}, \mathbf{q} - \mathbf{x} \rangle (\mathbf{u}(\mathbf{q}) - \mathbf{u}(\mathbf{x})) - \underline{\mathbb{K}}[\mathbf{p}] \langle \mathbf{x} - \mathbf{p}, \mathbf{q} - \mathbf{p} \rangle (\mathbf{u}(\mathbf{q}) - \mathbf{u}(\mathbf{p}))) dV_{\mathbf{q}} dV_{\mathbf{p}} + \mathbf{b}(\mathbf{x}) \\ &= \int \mathbf{C}(\mathbf{x}, \mathbf{q}) (\mathbf{u}(\mathbf{q}) - \mathbf{u}(\mathbf{x})) dV_{\mathbf{q}} + \mathbf{b}(\mathbf{x})\end{aligned}$$

where

$$\mathbf{C}(\mathbf{x}, \mathbf{q}) = \int (\underline{\mathbb{K}}[\mathbf{x}] \langle \mathbf{p} - \mathbf{x}, \mathbf{q} - \mathbf{x} \rangle - \underline{\mathbb{K}}[\mathbf{p}] \langle \mathbf{x} - \mathbf{p}, \mathbf{q} - \mathbf{p} \rangle + \underline{\mathbb{K}}[\mathbf{q}] \langle \mathbf{x} - \mathbf{q}, \mathbf{p} - \mathbf{q} \rangle) dV_{\mathbf{p}}$$

Equation of motion for a linearized material

- Equation of motion:

$$\rho \ddot{\mathbf{u}}(\mathbf{x}) = \int \mathbf{C}(\mathbf{x}, \mathbf{q})(\mathbf{u}(\mathbf{q}) - \mathbf{u}(\mathbf{x})) dV_{\mathbf{q}} + \mathbf{b}(\mathbf{x})$$

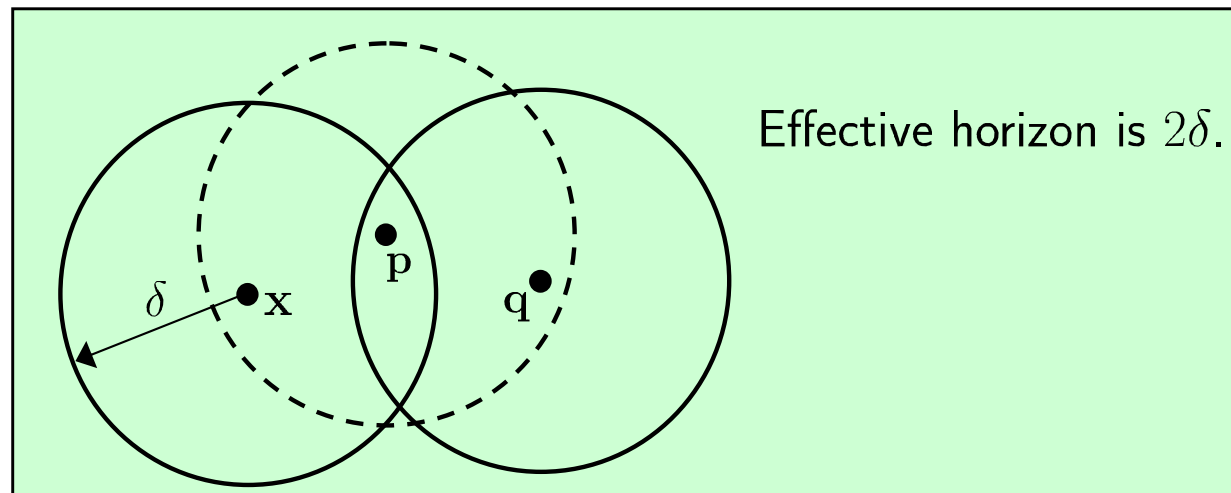
where

$$\mathbf{C}(\mathbf{x}, \mathbf{q}) = \int (\underline{\mathbb{K}}[\mathbf{x}]\langle \mathbf{p} - \mathbf{x}, \mathbf{q} - \mathbf{x} \rangle - \underline{\mathbb{K}}[\mathbf{p}]\langle \mathbf{x} - \mathbf{p}, \mathbf{q} - \mathbf{p} \rangle + \underline{\mathbb{K}}[\mathbf{q}]\langle \mathbf{x} - \mathbf{q}, \mathbf{p} - \mathbf{q} \rangle) dV_{\mathbf{p}}$$

- Similar structure to linear *bond-based* equation of motion but \mathbf{C} has different symmetry:

$$\mathbf{C}(\mathbf{q}, \mathbf{x}) = \mathbf{C}^T(\mathbf{x}, \mathbf{q}) \quad \dots \text{state-based}$$

$$\mathbf{C}(\mathbf{q}, \mathbf{x}) = \mathbf{C}(\mathbf{x}, \mathbf{q}) \quad \text{and} \quad \mathbf{C}(\mathbf{x}, \mathbf{q}) = \mathbf{C}^T(\mathbf{x}, \mathbf{q}) \quad \dots \text{bond-based}$$



Stability of a jump perturbation

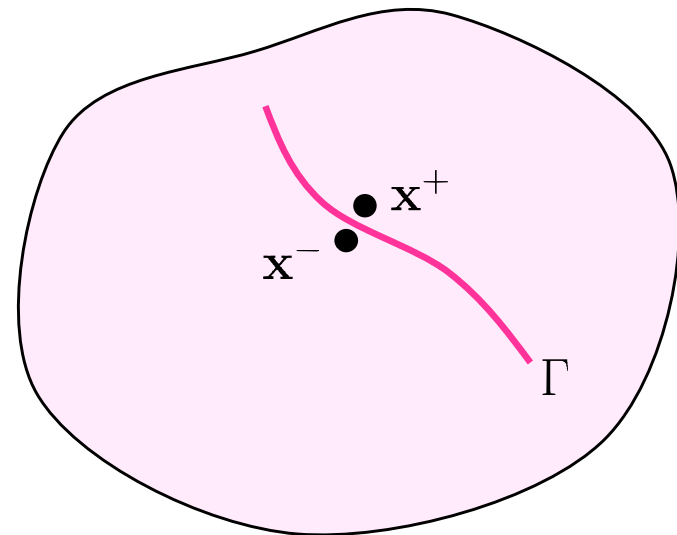
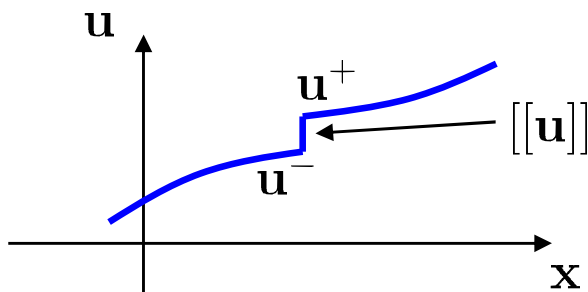
- Write the linearized equation of motion as:

$$\rho \ddot{\mathbf{u}}(\mathbf{x}) = \int \mathbf{C}(\mathbf{x}, \mathbf{q}) \mathbf{u}(\mathbf{q}) dV_{\mathbf{q}} - \mathbf{P}(\mathbf{x}) \mathbf{u}(\mathbf{x}) + \mathbf{b}(\mathbf{x})$$

where \mathbf{P} is the symmetric tensor defined by

$$\mathbf{P}(\mathbf{x}) = \int \mathbf{C}(\mathbf{x}, \mathbf{q}) dV_{\mathbf{q}} = \int \int \mathbb{K}[\mathbf{x}] \langle \mathbf{p} - \mathbf{x}, \mathbf{q} - \mathbf{x} \rangle dV_{\mathbf{p}} dV_{\mathbf{q}}$$

- Consider a small superposed displacement field \mathbf{u} containing a jump across a surface Γ .
- Define $[[\mathbf{u}]] = \mathbf{u}^+ - \mathbf{u}^-$.



Stability of a jump perturbation, ctd.

- Write the equation of motion on each side of the jump ($\mathbf{b} = \mathbf{0}$):

$$\rho \ddot{\mathbf{u}}^+ = \int \mathbf{C}(\mathbf{x}^+, \mathbf{q}) \mathbf{u}(\mathbf{q}) dV_{\mathbf{q}} - \mathbf{P}(\mathbf{x}^+) \mathbf{u}^+$$

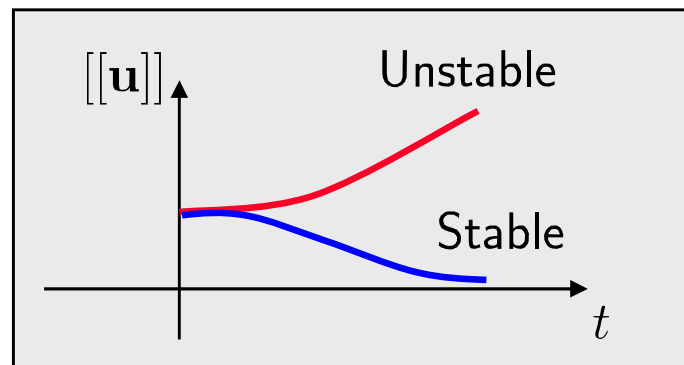
$$\rho \ddot{\mathbf{u}}^- = \int \mathbf{C}(\mathbf{x}^-, \mathbf{q}) \mathbf{u}(\mathbf{q}) dV_{\mathbf{q}} - \mathbf{P}(\mathbf{x}^-) \mathbf{u}^-$$

- \mathbf{C} and \mathbf{P} are continuous. Subtract.

$$\rho [[\ddot{\mathbf{u}}]] = -\mathbf{P} [[\mathbf{u}]]$$

$$\rho [[\ddot{\mathbf{u}}]] \cdot [[\mathbf{u}]] = -\mathbf{P} |[[\mathbf{u}]]|^2$$

- The jump grows if $[[\ddot{\mathbf{u}}]] \cdot [[\mathbf{u}]] > 0$. This can happen if \mathbf{P} has a negative eigenvalue.

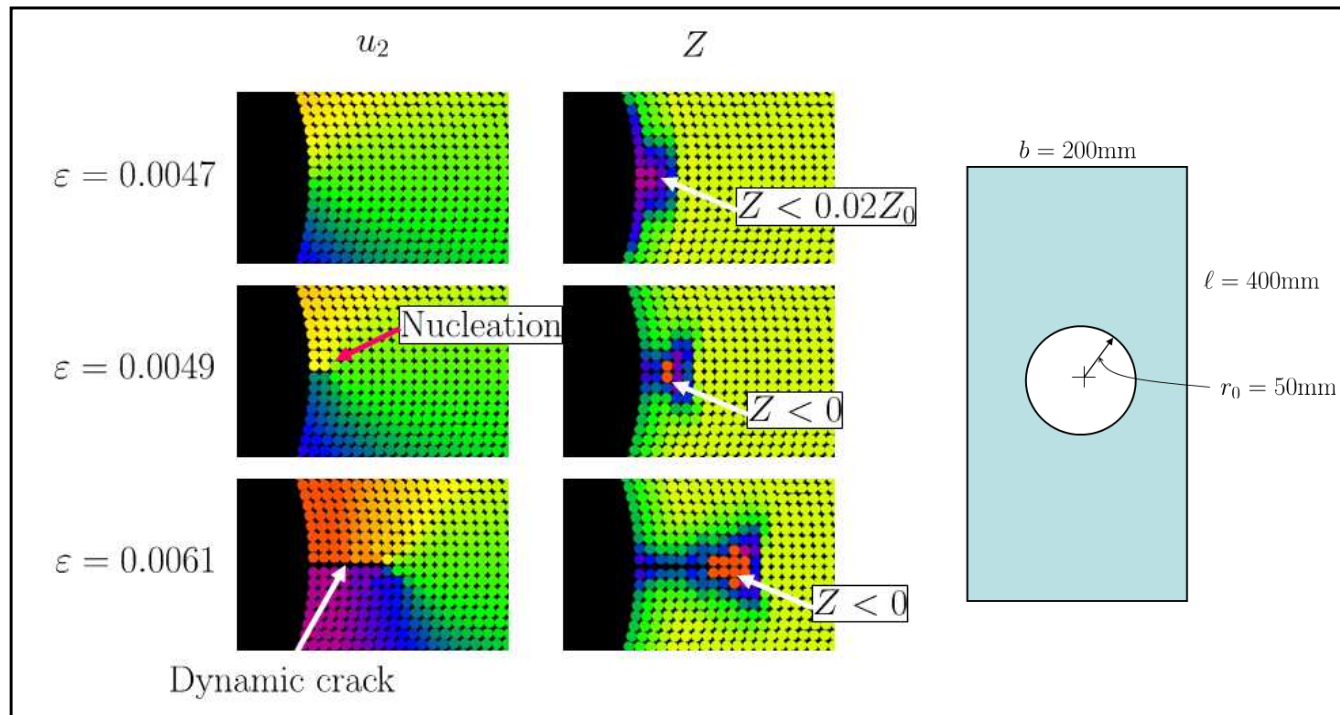


Crack nucleation condition

- Let the eigenvalues of $\mathbf{P}(\mathbf{x})$ be denoted $\{P_1, P_2, P_3\}$ and define the *stability index* by

$$Z(\mathbf{x}) = \min \{P_1, P_2, P_3\}.$$

- If $Z(\mathbf{x}) < 0$ then a crack can nucleate at \mathbf{x} .
- $Z(\mathbf{x})$ depends only on the material properties at \mathbf{x} .



Materials with a damage variable

- Assume there is a scalar *damage state* $\underline{\phi}$ such that

$$\psi = \psi(\underline{\mathbf{Y}}, \theta, \underline{\phi}) \quad \text{and} \quad \dot{\underline{\phi}} \geq 0. \quad \leftarrow \text{Damage is irreversible.}$$

- Repeat C-N argument to find that we still have (for $h = r = 0$)

$$\underline{\mathbf{T}} = \psi_{\underline{\mathbf{Y}}} \quad \text{and} \quad \eta = -\psi_{\theta}$$

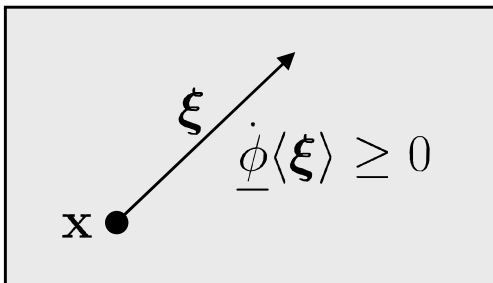
but now also have

$$\psi_{\underline{\phi}} \leq 0$$

and

$$\dot{\eta} = \frac{\dot{\psi}_d}{\theta} \quad \text{where} \quad \dot{\psi}_d := -\psi_{\underline{\phi}} \bullet \dot{\underline{\phi}}.$$

Energy dissipation





Damage evolution laws

- $\underline{\phi}\langle\xi\rangle$ is the damage in bond ξ (at some \mathbf{x}), determined by a damage evolution law:

$$\underline{\phi} = \underline{D}(\underline{\mathbf{Y}}, \underline{\dot{\mathbf{Y}}}, \dots)$$

- If $\underline{\mathbf{T}}\langle\xi\rangle = \mathbf{0}$ whenever $\underline{\phi}\langle\xi\rangle = 1$, the material has *strong* damage dependence (otherwise *weak*).



Damage evolution laws

Example: bond breakage

- Define the bond extension state by

$$\underline{e}\langle\xi\rangle = |\underline{\mathbf{Y}}\langle\xi\rangle| - |\xi|.$$

- Suppose

$$\underline{D}\langle\xi\rangle = H(\underline{e}_0\langle\xi\rangle, e_b)$$

where H =Heaviside step function and

$$\underline{e}_0\langle\xi\rangle = \max_t \langle\xi\rangle.$$

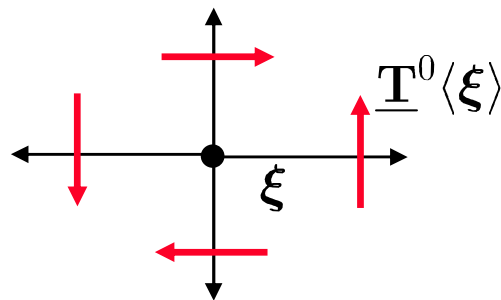
Damage in bond ξ jumps from 0 to 1 when its elongation exceeds the critical elongation e_b .

Damage in a constitutive model: have to be consistent with nonpolarity

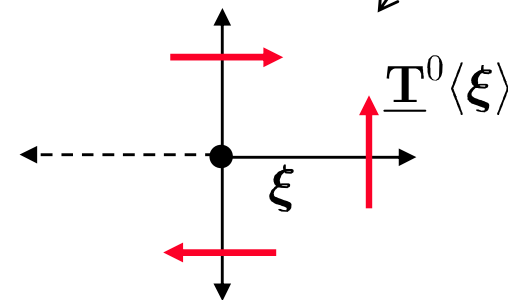
- Cannot in general do the following with a nonordinary material model $\underline{\mathbf{T}}^0$:

$$\underline{\mathbf{T}}\langle\xi\rangle = (1 - \phi\langle\xi\rangle)\underline{\mathbf{T}}^0\langle\xi\rangle$$

because the resulting model may fail to be nonpolar.



Four typical bonds in a
nonordinary material



Breaking a bond results in a net moment –
No longer nonpolar



Damage in a constitutive model: strong damage

- Suppose we have an elastic material with strain energy function $W = W^0(\underline{e})$. Then

$$\underline{\mathbf{T}}^0 = W_{\underline{\mathbf{Y}}} = W_{\underline{e}}^0 \underline{\mathbf{M}}, \quad \underline{\mathbf{M}} = \frac{\underline{\mathbf{Y}}}{|\underline{\mathbf{Y}}|}$$

- Define a material by

$$\psi(\underline{\mathbf{Y}}, \underline{\phi}) = W^0((1 - \underline{\phi})\underline{e}).$$

- Then

$$\underline{\mathbf{T}} = (1 - \underline{\phi})\underline{\mathbf{T}}^0$$

- Each bond has its force reduced by $1 - \underline{\phi}\langle \xi \rangle$.
- ψ is still objective so model is still nonpolar.



Damage in a constitutive model: separable damage

- Start with $W = W^0(\underline{\mathbf{Y}})$. Then

$$\underline{\mathbf{T}}^0 = W_{\underline{\mathbf{Y}}}^0$$

- Define a material by

$$\psi(\underline{\mathbf{Y}}, \underline{\phi}) = \Phi(\underline{\phi}) W^0(\underline{\mathbf{Y}})$$

where

$$\Phi(\underline{\phi}) = \frac{1}{V} \int_{\mathcal{H}} (1 - \underline{\phi}(\underline{\xi}))^2 dV_{\xi}.$$

- Find

$$\underline{\mathbf{T}} = \Phi(\underline{\phi}) \underline{\mathbf{T}}^0$$

- All the bond forces are multiplied by the same $\Phi(\underline{\phi})$.

Changing the length scale in a material model

- Suppose we want to change the horizon from δ_0 to δ_1 . Require the rescaled energy to be the same as the original W_0 if the deformation is homogeneous.
- The rescaled material model is

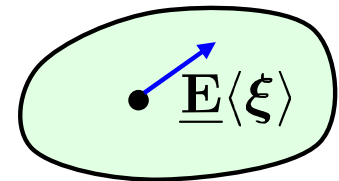
$$W_1(\underline{\mathbf{Y}}) = W_0(\underline{\mathbf{E}})$$

where $\underline{\mathbf{E}}$ is a state defined by

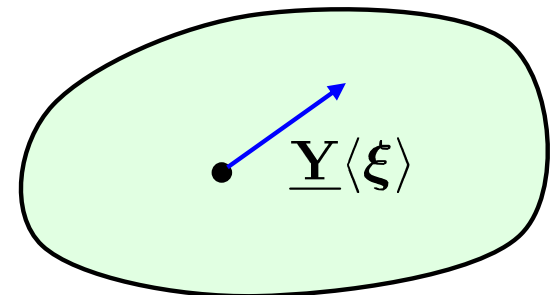
$$\underline{\mathbf{E}}\langle \xi \rangle = \frac{\delta_0}{\delta_1} \underline{\mathbf{Y}}\langle \xi \rangle$$

- Can show the force state scales according to

$$\underline{\mathbf{T}}_1(\underline{\mathbf{Y}}) = \left(\frac{\delta_1}{\delta_0} \right)^4 \underline{\mathbf{T}}_0(\underline{\mathbf{E}})$$



Original, δ_0

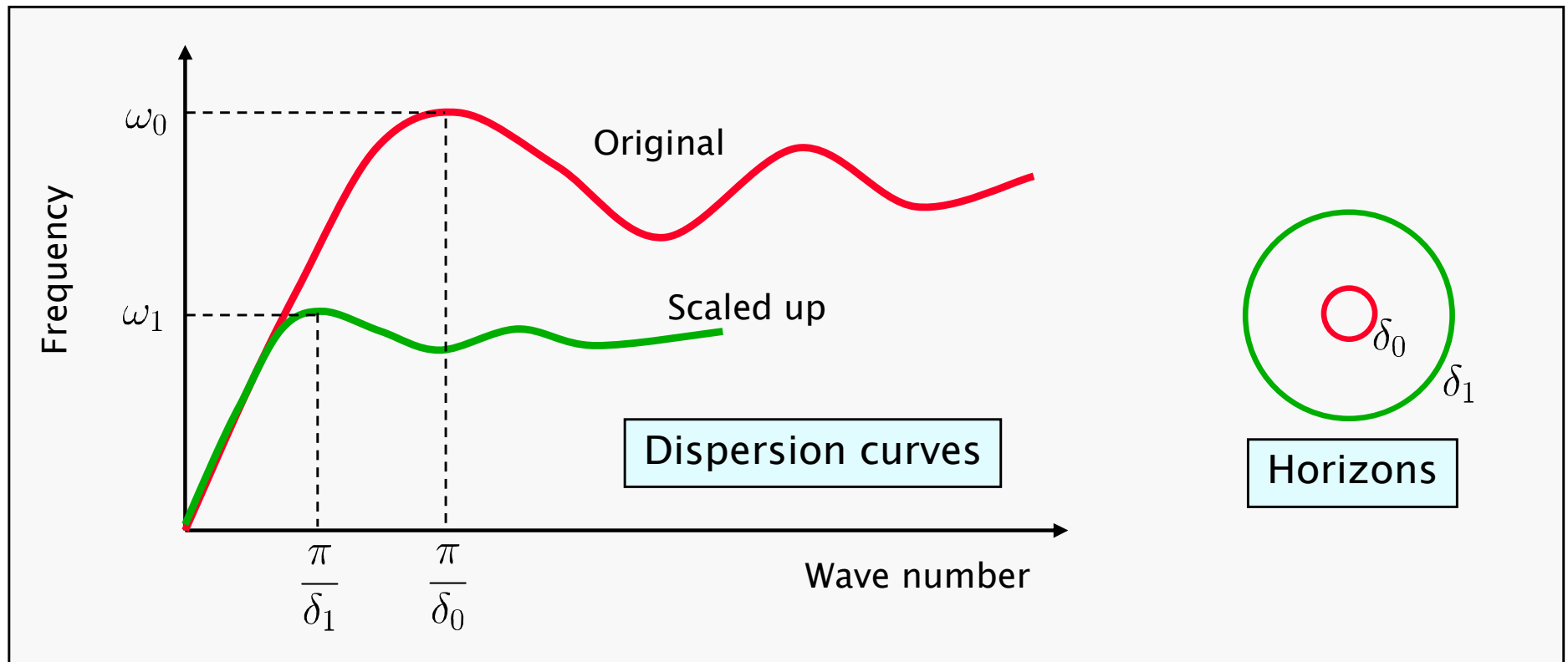


Rescaled, δ_1

Changing the length scale also changes the time scale

- Removing the small length scale also removes the high frequencies that characterize that length scale.

$$\delta_1 > \delta_0 \implies \omega_1 < \omega_0$$



Peridynamic stress tensor

In any peridynamic body, we can define a tensor field $\boldsymbol{\nu}$ such that:

- The force per unit area at \mathbf{x} through a plane with normal \mathbf{n} is

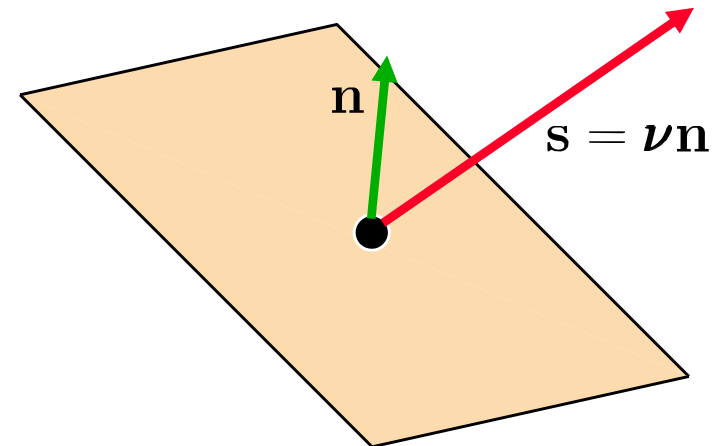
$$\mathbf{s} = \boldsymbol{\nu}(\mathbf{x})\mathbf{n}$$

- The peridynamic equation of motion can be written as

$$\rho \ddot{\mathbf{u}} = \text{div } \boldsymbol{\nu} + \mathbf{b}$$

i.e.,

$$\text{div } \boldsymbol{\nu}(\mathbf{x}) = \int \mathbf{f}(\mathbf{x}', \mathbf{x}) dV_{\mathbf{x}'}$$



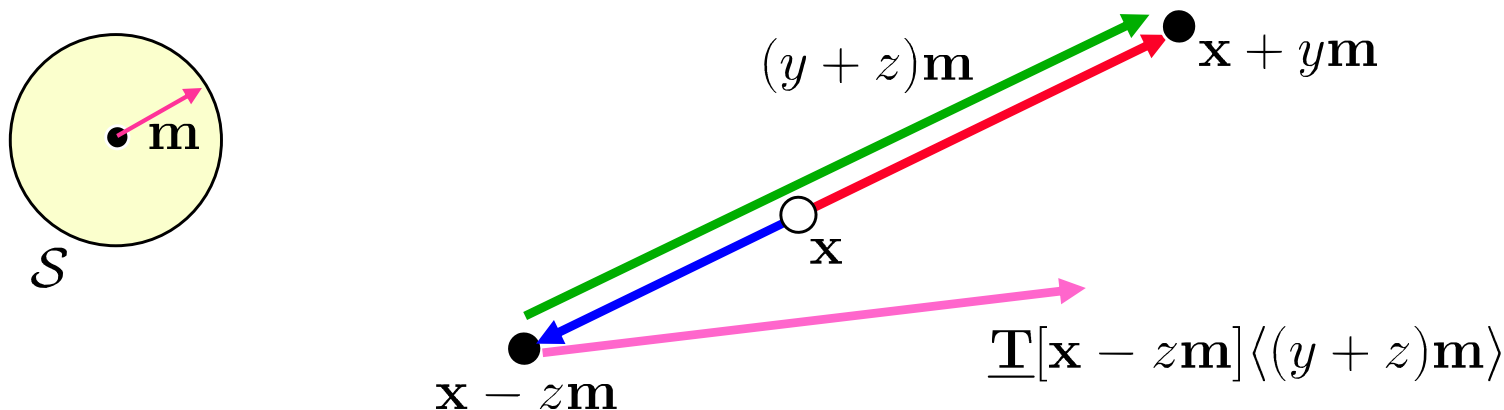
Peridynamic stress tensor, ctd.

- The peridynamic stress tensor is given by

$$\boldsymbol{\nu}(\mathbf{x}) = \int_{\mathcal{S}} \int_0^\infty \int_0^\infty (y+z)^2 \left((\underline{\mathbf{T}}[\mathbf{x}-z\mathbf{m}] \langle (y+z)\mathbf{m} \rangle) \otimes \mathbf{m} \right) dz dy d\Omega_{\mathbf{m}}$$

where \mathcal{S} is the unit sphere and Ω is solid angle.

- $\boldsymbol{\nu}$ sums up the forces in bonds that go through \mathbf{x} .



Convergence of peridynamics to the standard theory

Suppose the deformation is twice continuously differentiable. If the horizon is small, the deformation state is well approximated by

$$\underline{\mathbf{Y}}\langle\xi\rangle \approx (\nabla \mathbf{y})\xi$$

so we can write

$$W(\underline{\mathbf{Y}}) \approx W_c(\nabla \mathbf{y})$$

and it can be proven that

$$\boldsymbol{\nu} \approx \frac{\partial W_c}{\partial \nabla \mathbf{y}}$$

so $\boldsymbol{\nu}$ is basically a Piola-Kirchhoff stress tensor in a classical hyperelastic solid.

